



1 **Comparisons of stemflow yield and efficiency between two xerophytic**
2 **shrubs: the effects of leaves and implications in drought tolerance**

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13 **Abstract.**

14 Stemflow transports enriched precipitation to the rhizosphere and is highly important for
15 the survival of xerophytic shrubs in water-stressed ecosystems. However, its ecological
16 significance has generally been underestimated because it is relatively limited in amount, and
17 the biotic mechanisms that affect it have not been thoroughly studied at the leaf scale. In this
18 study, the branch stemflow volume (SF_b), the shrub stemflow equivalent water depth (SF_d),
19 the stemflow percentage of incident precipitation (SF%), the stemflow productivity (SFP),
20 the funnelling ratio (FR), the rainfall characteristics and the plant traits of branches and
21 leaves of *C. korshinskii* and *S. psammophila* were measured during the 2014 and 2015 rainy
22 seasons in the northern Loess Plateau of China. This study evaluated the stemflow production
23 efficiency for the first time with the combined results of SFP and FR, and sought to determine
24 the inter- and intra-specific differences in stemflow production and production efficiency, as
25 well as the specific bio-/abiotic mechanisms that affected stemflow. The results indicated that
26 precipitation amount was the most influential rainfall characteristic that affected stemflow in
27 these two endemic shrub species and that stem biomass and leaf biomass were the most
28 influential plant traits in *C. korshinskii* and *S. psammophila*, respectively. *C. korshinskii* had a
29 greater stemflow production and production efficiency at all precipitation levels, and the
30 largest inter-specific difference was generally in the 5–10-mm young shoots during the most
31 frequent rainfall events of ≤ 2 mm. *C. korshinskii* had a lower precipitation threshold (0.9 mm
32 vs. 2.1 mm for *S. psammophila*), which provided more available water from rainfall for
33 stemflow. The leaves affected stemflow production, and the beneficial leaf traits contributed
34 to the higher stemflow production of *C. korshinskii*. In summary, *C. korshinskii* might have
35 greater drought tolerance and a competitive edge in a dryland ecosystem because of greater
36 and more efficient stemflow production, a lower precipitation threshold and more
37 advantageous leaf traits.

38 **Keywords:** Xerophytic shrub; Stemflow production; stemflow production efficiency;
39 Threshold precipitation; Beneficial leaf traits.



40 **1 Introduction**

41 Stemflow channels divert precipitation pointedly into the root zone of a plant via
42 preferential root paths, worm paths and soil macropores. The double-funnelling effects of
43 stemflow and preferential flow create “hot spots” and “hot moments” by enhancing
44 biogeochemical reactivity at the terrestrial-aquatic interface (McClain et al., 2003; Johnson
45 and Lehmann, 2006), thus substantially contributing to the formation and maintenance of
46 so-called “fertile islands” (Whitford et al., 1997), “resource islands” (Reynolds et al., 1999)
47 or “hydrologic islands” (Rango et al., 2006). This effect is important for the normal function
48 of rain-fed dryland ecosystems (Wang et al., 2011).

49 Shrubs are a representative plant functional type (PFT) in dryland ecosystems and have
50 developed effective physiological drought tolerance by reducing water loss, e.g., through
51 adjusting their photosynthetic and transpiration rate by regulating stomatal conductance and
52 abscisic acid (ABA), titling their osmotic equilibrium by regulating the concentration of
53 soluble sugars and inorganic ions, and removing free radicals (Ma et al., 2004, 2008). The
54 efficient production of stemflow is a vital eco-hydrological flux involved in soil water
55 replenishment (Pressland 1973) as well as an effective strategy to acquire water (Murakami,
56 2009) and withstand drought (Martinez-Meza and Whitford, 1996). However, because
57 stemflow occurs in small amounts, previous studies have usually ignored stemflow (Llorens
58 and Domingo, 2007) and have underestimated its disproportionately high influence on the
59 survival and competitiveness of xerophytic shrub species. Therefore, the quantification of
60 inter- and intra-specific stemflow production is important to assess the stemflow production
61 efficiency and to elucidate the underlying bio-/abiotic mechanisms.

62 Stemflow production includes the stemflow volume and depth, and it describes the total
63 flux channelled down to the base of a branch or a trunk, but stemflow data are unavailable for
64 comparison of inter-specific differences caused by variations in the branch architecture, the



65 canopy structure, the shrub species and the eco-zone. Herwitz (1986) introduced the
66 funnelling ratio (FR), which is expressed as the quotient of the volume of stemflow produced
67 and the product of the base area and the precipitation amount. It indicates the efficiency with
68 which individual branches or shrubs capture raindrops and deliver the water to the root zone
69 (Siegert and Levia, 2014). The FR allows a comparison of the inter- and intra-specific
70 stemflow production under different precipitation conditions. However, the FR does not
71 provide a connection between hydrological processes (e.g., rainfall redistribution) and the
72 plant growth processes (e.g., biomass accumulation and allocation). Recently, Yuan et al.
73 (2016) have introduced the parameter stemflow productivity (SFP), expressed as the volume
74 of stemflow production per unit of branch biomass. The SFP describes the efficiency in an
75 energy-conservation manner by comparing the stemflow volume of a unit biomass increment
76 of different-sized branches.

77 The precipitation amount is an abiotic mechanism that has been recognized as the single
78 most influential rainfall characteristic (Clements 1972; André et al., 2008; Van Stan et al.,
79 2014). However, in terms of biotic mechanisms, although the canopy structure (Mauchamp
80 and Janeau, 1993; Crockford and Richardson, 2000; Pypker et al., 2011) and branch
81 architecture (Herwitz, 1987; Murakami 2009; (Herwitz, 1987;Murakami, 2009;
82 Carlyle-Moses and Schooling, 2015) have been studied for years, the most important plant
83 traits that vary with location and shrub species have not yet been determined. The effects of
84 the leaves have been studied more recently at a smaller scale, e.g., leaf orientation (Crockford
85 and Richardson, 2000), shape (Xu et al., 2005), arrangement pattern (Owens et al., 2006),
86 pubescence (Garcia-Estringana et al., 2010), area (Sellin et al., 2012), epidermis microrelief
87 (Roth-Nebelsick et al., 2012), amount (Li and Xiao, 2016), biomass (Yuan et al., 2016; Li et
88 al., 2016), etc. Although comparisons of stemflow production during the foliated and dormant
89 seasons usually indicate negative effects of leaves because the more stemflow occurred at the



90 leafless period (Dolman, 1987; Neal et al., 1993; Muzyło et al., 2012), both negligible and
91 positive effects have also been confirmed by Martinez-Meza and Whitford (1996) and Liang
92 et al. (2009), respectively. Nevertheless, the validity of these findings has been called into
93 question as a result of the seasonal variation of meteorological conditions and plant traits, e.g.,
94 wind speed (André et al., 2008), rainfall intensity (Dunkerley et al., 2014 a, b), air
95 temperature and consequent precipitation type (snow-to-rain vs. snow) (Levia, 2004).
96 Therefore, a controlled experiment with foliated and manually defoliated plants under the
97 same stand conditions is needed to resolve these uncertainties.

98 In this study, the branch stemflow volume (SF_b), the shrub stemflow depth (SF_d), the
99 stemflow percentage of the incident precipitation amount (SF%), the SFP and the FR were
100 measured in two shrub species (*C. korshinskii* and *S. psammophila*) endemic to a semiarid
101 area of northern China during the 2014 and 2015 rainy seasons. The objectives of this study
102 were to (1) quantify the inter- and intra-specific stemflow production (SF_b , SF_d and SF%) and
103 the production efficiency (SFP and FR); (2) investigate the effects of the rainfall
104 characteristics and plant traits on the stemflow in these two shrub species; and (3) specifically
105 identify leaf characteristics that affect the stemflow with respect to morphology, structural
106 characteristics and the biomass partitioning pattern. The achievement of these research
107 objectives would provide a novel characterization of plant drought tolerance and species
108 competitiveness in terms of stemflow and further the understanding of the effects of leaves on
109 the survival and growth of plants from an eco-hydrological perspective.

110

111 **2 Materials and Methods**

112 **2.1 Study area**

113 This study was conducted at the Liudaogou catchment (110°21'-110°23'E,
114 38°46'-38°51'N) in Shenmu County in the Shaanxi Province of China. It is 6.89 km² and



115 1094-1273 m above sea level (a.s.l.). This area has a semiarid continental climate with
116 well-defined rainy and dry seasons. The mean annual precipitation (MAP) between 1971 and
117 2013 was 414 mm, with approximately 77% of the annual precipitation amount occurring
118 during the rainy season (Jia et al., 2013), which lasts from July to September. The mean
119 annual temperature and potential evaporation are 9.0 °C and 1337 mm year⁻¹ (Zhao and Shao,
120 2009), respectively. The coldest and warmest months are January and July, with an average
121 monthly temperature of 9.7 °C and 23.7 °C, respectively. Two soil types of Aeolian sandy soil
122 and Ust-Sandiic Entisol dominate this catchment (Jia et al., 2011). Soil particles consist of
123 11.2%-14.3% clay, 30.1%-44.5% silt and 45.4%-50.9% sand in terms of the soil classification
124 system of United States Department of Agriculture (Zhu and Shao, 2008). The original plants
125 are scarcely present, except for very few surviving shrub species, e.g., *Ulmus macrocarpa*,
126 *Xanthoceras sorbifolia*, *Rosa xanthina*, *Spiraea salicifolia*, etc. The currently predominant
127 shrub species were planted decades ago, e.g., *S. psammophila*, *C. Korshinskii*, *Amorpha*
128 *fruticosa*, etc., and the predominant grass species include *Medicago sativa*, *Stipa bungeana*,
129 *Artemisia capillaris*, *Artemisia sacrorum*, etc. (Ai et al., 2015).

130 *C. Korshinskii* and *S. psammophila* are endemic shrub species in arid and semiarid
131 northern China and were planted for wind-proofing and dune-stabilizing because of their
132 great drought tolerance. Two representative experimental stands were established in the
133 southwest of the Liudaogou catchment (Fig. 1). Both *C. korshinskii* and *S. psammophila* were
134 planted approximately twenty years ago, and the two stands share a similar slope of 13-18 °, a
135 size of 3294-4056 m², and an elevation of 1179-1207 m a.s.l. However, the *C. korshinskii*
136 experimental stand had a 224 ° aspect with a loess ground surface, whereas the *S.*
137 *psammophila* experimental stand had a 113 ° aspect with a sand ground surface.

138

139 Fig. 1. Location of the experimental stands and facilities for stemflow measurements of *C.*
140 *korshinskii* and *S. psammophila* at the Liudaogou catchment in the Loess Plateau of China.

141



142 2.2 Field experiments

143 Field experiments were conducted during the rainy seasons of 2014 (July 1 to October 3)
144 and 2015 (June 1 to September 30) to measure the rainfall characteristics, plant traits and
145 stemflow. To avoid the effects of gully micro-geomorphology on recording the rainfall
146 characteristics, we installed an Onset® (Onset Computer Corp., Bourne, MA, USA) RG3-M
147 tipping bucket rain gauge (0.2 mm per tip) at each experimental stand. Three 20-cm-diameter
148 rain gauges were placed around to adjust the inherent underestimating of automatic
149 precipitation recording (Groisman and Legates, 1994). Then, rainfall duration (RD, h),
150 rainfall interval (RI, h), the average rainfall intensity (I, mm h⁻¹), the maximum rainfall
151 intensity in 5 min (I₅, mm h⁻¹), 10 min (I₁₀, mm h⁻¹) and 30 min (I₃₀, mm h⁻¹) could be
152 calculated accordingly. In this study, the individual rainfall events were greater than 0.2 mm
153 and separated by a period of at least four hours without rain (Giacomin and Trucchi, 1992).

154 *C. korshinskii* and *S. psammophila*, as modular organisms and multi-stemmed shrub
155 species, have branches of that exist as independent individuals. Therefore, we focused on the
156 inter- and intra-specific branch stemflow by experimenting on sample shrubs that had a
157 similar canopy structure. Four mature shrubs were selected for *C. korshinskii* (designated as
158 C1, C2, C3 and C4) and *S. psammophila* (designated as S1, S2, S3 and S4) for the stemflow
159 measurements. They had isolated canopies, similar intra-specific heights and canopy areas,
160 e.g., 2.1 ± 0.2 m and 5.14 ± 0.26 m² for C1-C4, and 3.5 ± 0.2 m and 21.35 ± 5.21 m² for
161 S1-S4. We measured the morphological characteristics of all the 180 branches of C1-C4 and
162 all the 261 branches of S1-S4, including the branch basal diameter (BD, mm), branch length
163 (BL, cm) and branch inclination angle (BA, °). The leaf area index (LAI) and the foliage
164 orientation (MTA, the mean tilt angle of leaves) were measured using LiCor® (LiCor
165 Biosciences Inc., Lincoln, NE, USA) 2200C plant canopy analyser approximately twice a
166 month.



167 A total of 53 branches of *C. korshinskii* and 98 branches of *S. psammophila* were
168 selected for stemflow measurements following the criteria: 1) no intercrossing stems; 2) no
169 turning point in height from branch tip to the base; 3) representativeness in amount and
170 branch size. Stemflow was collected using aluminum foil collars, which was fitted around the
171 entire branch circumference and sealed by neutral silicone caulking (Fig. 1). A
172 0.5-cm-diameter PVC hose led the stemflow to lidded containers. The stemflow volume was
173 measured within two hours after the rainfall ended during the daytime; if the rainfall ended at
174 night, we took the measurement early the next morning.

175 Another three shrubs of each species were destructively measured for biomass and leaf
176 traits. They had similar canopy heights and areas as those of the shrubs for which the
177 stemflow was measured and were designated as C5-C7 (2.0-2.1 m and 5.84-6.77 m²) and
178 S5-S7 (3.0-3.4 m and 15.43-19.20 m²), thus allowing the development of allometric models
179 for the estimation of the corresponding biomass and leaf traits of C1-C4 and S1-S4 (Levia
180 and Herwitz, 2005; Siles et al., 2010a, 2010b; Stephenson et al., 2014). A total of 66 branches
181 for C5-C7 and 61 branches for S5-S7 were measured when the shrubs showed maximum
182 vegetative growth during mid-August for the biomass of leaves and stems (BML and BMS,
183 g), the leaf area of the branches (LAB, cm²), and the leaf numbers of the branches (LNB).
184 The BML and BMS were weighted after oven-drying of 48 hours. The detailed measurements
185 have been reported in Yuan et al., (2016). The validity of the allometric models was verified
186 by measuring another 13 branches of C5-C7 and 14 branches of S5-S7.

187

188 2.3 Calculations

189 Biomass and leaf traits were estimated by allometric models as an exponential function
190 of BD (Siles et al., 2010a, b; Jonard et al., 2006):

$$191 \quad PT_c = a * BD^b \quad (1)$$



192 where a and b are constants, and PT_e refers to the estimated plant traits BML, BMS, LAB
 193 and LNB. The other plant traits could be calculated accordingly, including individual leaf
 194 area of branch ($ILAB = 100 \cdot LAB / LNB$, mm^2), the percentage of stem biomass to that of
 195 branch ($PBMS = BMS / (BML + BMS) \cdot 100\%$, %), specific leaf weight ($SLW = BML / LAB$,
 196 g cm^{-2}), Huber value ($HV = BBA / LAB = 3.14 \cdot BD^2 / (400 \cdot LAB)$, unitless, where BBA is the
 197 branch basal area (cm^2)).

198 In this study, stemflow production was defined as the branch volume production
 199 (hereafter “stemflow production”, SF_b , mL), the equivalent water depth on the basis of shrub
 200 canopy area (hereafter “stemflow depth”, SF_d , mm), and the stemflow percentage of the
 201 incident precipitation amount (hereafter “stemflow percentage”, SF%, %):

$$202 \quad SF_d = 10 \cdot \sum_{i=1}^n SF_{bi} / CA \quad (2)$$

$$203 \quad SF\% = (SF_d / P) \cdot 100\% \quad (3)$$

204 where SF_{bi} is the volume of stemflow production of branch i (mL), CA is the canopy area
 205 (cm^2), n is the number of branches, and P is the incident precipitation amount (mm).

206 Stemflow productivity (SFP, mL g^{-1}) was expressed as the SF_b (mL) of unit branch
 207 biomass (g) and represented the stemflow production efficiency of different-sized branches in
 208 terms of energy-conservation:

$$209 \quad SFP = SF_b / (BML + BMS) \quad (4)$$

210 The funnelling ratio (FR) was computed as the quotient of SF_b and the product of P and
 211 BBA (Herwitz, 1986). A FR with a value greater than 1 indicated a positive effect of the
 212 canopy on the stemflow production (Carlyle-Moses and Price, 2006). The value of (P * BBA)
 213 equals to the precipitation amount that would have been caught by the rain gauge occupying
 214 the same basal area at the clearing:

$$215 \quad FR = 10 \cdot SF_b / (P \cdot BBA) \quad (5)$$

216



217 **2.4 Data analysis**

218 A Pearson correlation analysis was performed to test the relationship between SF_b and
219 each of the rainfall characteristics and plant traits. Significantly correlated variables were
220 further tested with a partial correlation analysis for their separate effects on SF_b . Then, the
221 qualified variables were fed into a stepwise regression with forward selection to identify the
222 most influential bio-/abiotic factors (Carlyle-Moses and Schooling, 2015; Yuan et al., 2016).
223 Similarly to a principal component analysis and ridge regression, stepwise regression has
224 commonly been used because it gets a limited effect of multicollinearity (Návar and Bryan,
225 1990; Honda et al., 2015; Carlyle-Moses and Schooling, 2015). Moreover, we excluded
226 variables that had a variance inflation factor (VIF) greater than 10 to minimize the effects of
227 multicollinearity (O'Brien, 2007). The same analysis method was also applied to identify the
228 most influential bio-/abiotic factors affecting SFP and FR. The level of significance was set at
229 95% confidence interval ($p = 0.05$). The SPSS 20.0 (IBM Corporation, Armonk, NY, USA),
230 Origin 8.5 (OriginLab Corporation, Northampton, MA, USA), and Excel 2013 (Microsoft
231 Corporation, Redmond, WA, USA) were used for data analysis.

232

233 **3 Results**

234 **3.1 Species-specific variation of plant traits**

235 According to the *Flora of China* and the field observation, both *C. korshinskii* and *S.*
236 *psammophila* had an inverted-cone canopy and no trunk, with the branches running obliquely
237 from the base. *S. psammophila* usually grew to 3-4 m and had an odd number of strip-shaped
238 leaves of 24-mm in width and 4080-mm in length. The young leaves were pubescent and
239 gradually became subglabrous (Chao and Gong, 1999) (Fig. 2). In comparison, *C. korshinskii*
240 usually grew to 2 m and had pinnate compound leaves with 12-16 foliates in an opposite or
241 sub-opposite arrangement (Wang et al., 2013). The leaf was concave and lanceolate-shaped,



242 with an acute leaf apex and an obtuse base. Both sides of the leaves were densely sericeous
243 with appressed hairs (Liu et al., 2010) (Fig. 2).

244

245 Fig. 2. Comparison of leaf morphologies of *C. korshinskii* and *S. psammophila*.
246

247 Allometric models were developed to estimate the biomass and leaf traits of the
248 branches of *C. korshinskii* and *S. psammophila* measured for stemflow. The quality of the
249 estimates was verified by linear regression. As shown in Fig. 3, the regression of LAB, LNB,
250 BML and BMS of *C. korshinskii* had an approximately 1:1 slope (0.99 for the biomass
251 indicators and 1.04 for the leaf traits) and an R^2 value of 0.93-0.95. According to Yuan et al.,
252 (2016), the regression of *S. psammophila* had a slope of 1.13 and an R^2 of 0.92. Therefore,
253 those allometric models were appropriate.

254

255 Fig. 3. Verification of the allometric models for estimating the biomass and leaf traits of *C.*
256 *korshinskii*. BML and BMS refer to the biomass of the leaves and stems, respectively, and
257 LAB and LNB refer to the leaf area and the number of branches, respectively.
258

259 *C. korshinskii* had a similar average branch size and angle, but a shorter branch length
260 than did *S. psammophila*, e.g., 12.48 ± 4.16 mm vs. 13.73 ± 4.36 mm, $60 \pm 18^\circ$ vs. $60 \pm 20^\circ$,
261 and 161 ± 35 cm vs. 267.3 ± 49.7 cm, respectively. Regarding branch biomass accumulation,
262 *C. korshinskii* had a smaller BML (an average of 19.93 ± 10.81 g) and a larger BMS (an
263 average 141.07 ± 110.78 g) than did *S. psammophila* (an average of 27.85 ± 20.71 g and
264 130.65 ± 101.35 g, respectively). Both the BML and BMS increased with increasing branch
265 size for these two shrub species. When expressed as a proportion, *C. korshinskii* had a larger
266 PBMS than that of *S. psammophila* in all the BD categories. The PBMS-specific difference
267 increased with an increasing branch size, ranging from 1.24% for the 5–10-mm branches to
268 7.22% for the >18-mm branches.

269 Although an increase in LAB and LNB and a decrease in ILAB were observed for both



270 shrub species with an increase in branch size, *C. korshinskii* had a larger LAB (an average of
 271 $2509.05 \pm 1355.30 \text{ cm}^2$) and LNB (an average of 12479 ± 8409), but a smaller ILAB (an
 272 average of $21.94 \pm 2.99 \text{ mm}^2$) than did *S. psammophila* for each BD level (Table 1). The
 273 inter-specific differences in the leaf traits decreased with increasing branch size. The largest
 274 difference occurred for the 5–10-mm branches, e.g., LNB and LAB were 12.21-fold and
 275 2.41-fold larger for *C. korshinskii*, and ILAB was 5.32-fold larger for *S. psammophila*. *C.*
 276 *korshinskii* had a larger SLW (an average of $126.04 \pm 0.29 \text{ g cm}^{-2}$) and HV (0.0507 ± 0.0064)
 277 than did *S. psammophila* ($73.87 \pm 14.52 \text{ g cm}^{-2}$ and 0.0009 ± 0.0001 , respectively). As the
 278 branch size increased, the SLW of *S. psammophila* decreased from 95.62 g cm^{-2} for the 5–
 279 10-mm branches to 58.07 g cm^{-2} for the >18-mm branches, but the HV of *C. korshinskii*
 280 increased from 0.0438 to 0.0615.

281
 282 Table 1. Comparison of branch morphology, biomass and leaf traits of *C. korshinskii* and *S.*
 283 *psammophila*.
 284

285 3.2 Stemflow production of *C. korshinskii* and *S. psammophila*

286 In this study, stemflow production was expressed as SF_b on the branch scale and SF_d and
 287 $SF\%$ on the shrub scale. The SF_b was an average of 290.6 mL and 150.3 mL for individual
 288 branches of *C. korshinskii* and *S. psammophila*, respectively. The SF_b was positively
 289 correlated with the branch size and precipitation of these two shrub species. As the branch
 290 size increased, SF_b increased from 119.0 mL for the 5–10-mm branches to 679.9 mL for
 291 the >20-mm branches for *C. korshinskii* and from 43.0 mL to 281.8 mL for the corresponding
 292 BD categories of *S. psammophila*. However, with increasing precipitation, a larger
 293 intra-specific difference in SF_b was observed, which increased from 28.4 mL during rains ≤ 2
 294 mm to 771.4 mL during rains >20 mm for *C. korshinskii* and from 9.0 mL to 444.3 mL for
 295 the corresponding precipitation categories of *S. psammophila*. The intra-specific differences
 296 in SF_b were significantly affected by the rainfall characteristics and the plant traits. Up to



297 2375.9 mL of stemflow was measured for the >18-mm branches of *C. korshinskii* during
298 rains >20 mm, but only 6.8 mL of stemflow occurred for the 5–10-mm branches during rains
299 ≤ 2 mm. For comparison, a maximum SF_b of 2097.6 mL and a minimum of 1.8 mL were
300 measured for *S. psammophila*.

301 *C. korshinskii* produced a larger SF_b than did *S. psammophila* for all BD and
302 precipitation categories, and the inter-specific differences in SF_b also varied substantially
303 with the rainfall characteristics and the plant traits. A maximum difference of 4.3-fold larger
304 for the SF_b of *C. korshinskii* was observed for the >18-mm branches during rains ≤ 2 mm. As
305 the precipitation increased, the SF_b -specific difference decreased from 3.2-fold larger for *C.*
306 *korshinskii* during rains ≤ 2 mm to 1.7-fold larger during rains >20 mm. The largest
307 SF_b -specific difference occurred for the 5–10-mm branches for almost all precipitation
308 categories, but no clear trend of change was observed with increasing branch size (Table 2).

309 SF_d and SF% averaged 1.00 mm and 8.0%, respectively, for individual *C. korshinskii*
310 shrubs and 0.8 mm and 5.5%, respectively, for individual *S. psammophila* shrubs. These
311 parameters increased with increasing precipitation, ranging from 0.09 mm and 5.8% during
312 rains ≤ 2 mm to 2.64 mm and 8.9% during rains >20 mm for *C. korshinskii* and from 0.01 mm
313 and 0.7% to 2.23 mm and 7.9% for the corresponding precipitation categories of *S.*
314 *psammophila*, respectively. Additionally, the individual *C. korshinskii* shrubs had a larger
315 stemflow than did *S. psammophila* for all precipitation categories. The maximum differences
316 in SF_d and SF% were 8.5- and 8.3-fold larger for *C. korshinskii* during rains ≤ 2 mm and
317 decreased with increasing precipitation to 1.2- and 1.1-fold larger during rains >20 mm.

318
319 Table 2. Comparison of stemflow production (SF_b , SF_d and SF%) between *C. korshinskii* and
320 *S. psammophila*.
321

322 3.3 Stemflow production efficiency of *C. korshinskii* and *S. psammophila*

323 Combined results for SFP and FR, the stemflow production efficiency were assessed for



324 *C. korshinskii* and *S. psammophila*. SFP averaged 1.95 mL g⁻¹ and 1.19 mL g⁻¹ for individual
325 *C. korshinskii* and *S. psammophila* branches, respectively (Table 3). As precipitation
326 increased, SFP increased from 0.19 mL g⁻¹ during rains ≤2 mm to 5.08 mL g⁻¹ during
327 rains >20 mm for *C. korshinskii* and from 0.07 mL g⁻¹ to 3.43 mL g⁻¹ for the corresponding
328 precipitation categories for *S. psammophila*. With an increase in branch size, SFP decreased
329 from 2.19 mL g⁻¹ for the 5–10-mm branches to 1.62 mL g⁻¹ for the >18-mm branches of *C.*
330 *korshinskii* and from 1.64 mL g⁻¹ to 0.80 mL g⁻¹ for the corresponding BD categories of *S.*
331 *psammophila*. Maximum SFP values of 5.60 mL g⁻¹ and 4.59 mL g⁻¹ were recorded for *C.*
332 *korshinskii* and *S. psammophila*, respectively. Additionally, *C. korshinskii* had a larger SFP
333 than that of *S. psammophila* for all precipitation and BD categories. This inter-specific
334 difference in SFP decreased with increasing precipitation from 2.5-fold larger for *C.*
335 *korshinskii* during rains ≤2 mm to 1.5-fold larger during rains >20 mm, and it increased with
336 increasing branch size: from 1.3-fold larger for *C. korshinskii* for the 5–10-mm branches to
337 2.0-fold larger for the >18-mm branches.

338
339 Table 3. Comparison of stemflow productivity (SFP) between *C. korshinskii* and *S.*
340 *psammophila*.
341

342 FR averaged 172.3 and 69.3 for the individual branches of *C. korshinskii* and *S.*
343 *psammophila*, respectively (Table 4). As the precipitation increased, an increasing trend was
344 observed, ranging from 129.2 during rains ≤2 mm to 190.3 during rains >20 mm for *C.*
345 *korshinskii* and from 36.7 to 96.1 during the corresponding precipitation categories for *S.*
346 *psammophila*. FR increased with increasing BA from 149.9 for the ≤30°-branches to 198.2
347 for the >80°-branches of *C. korshinskii* and from 55.0 to 85.6 for the corresponding BA
348 categories of *S. psammophila*. Maximum FR values of 276.0 and 115.7 were recorded for *C.*
349 *korshinskii* and *S. psammophila*, respectively. Additionally, *C. korshinskii* had a larger FR
350 than *S. psammophila* for all precipitation and BA categories. The inter-specific difference in



351 FR decreased with increasing precipitation from the 3.5-fold larger for *C. korshinskii* during
352 rains ≤ 2 mm to 2.0-fold larger during rains > 20 mm, and it decreased with an increase in the
353 branch inclination angle: from 2.7-fold larger for *C. korshinskii* for the $\leq 30^\circ$ -branches to
354 2.3-fold larger for the $> 80^\circ$ -branches.

355

356 Table 4. Comparison of the funnelling ratio (FR) for *C. korshinskii* and *S. psammophila*.

357

358 **3.4 Bio/abiotic influential factors of stemflow production and production efficiency**

359 For both *C. korshinskii* and *S. psammophila*, BA was the only plant trait that had no
360 significant correlation with SF_b ($r < 0.13$, $p > 0.05$) as indicated by Pearson correlation
361 analysis. The separate effects of the remaining plant traits were verified by using a partial
362 correlation analysis, but BL, ILAB and PBMS failed this test. The remaining plant traits,
363 including BD, LAB, LNB, BML and BMS, were regressed with SF_b by using the forward
364 selection method. Biomass was finally identified as the most important biotic indicator that
365 affected stemflow, which behaved differently in *C. korshinskii* for BMS and in *S.*
366 *psammophila* for BML. The same analysis methods indicated that the precipitation amount
367 was the most important rainfall characteristic that affected stemflow in these two shrub
368 species.

369 SF_b and SF_d had a good linear relationship with the precipitation amount ($R^2 \geq 0.93$) for
370 both shrub species (Fig. 4). The > 0.9 -mm and > 2.1 -mm rains were required to start SF_b for *C.*
371 *korshinskii* and *S. psammophila*, respectively, results consistent with the 0.8-mm and 2.0-mm
372 precipitation threshold calculated with SF_d . Moreover, the precipitation threshold increased
373 with increasing branch size. The precipitation threshold values were 0.69 mm, 0.72 mm, 1.35
374 mm and 0.81 mm for the 5–10-mm, 10–15-mm, 15–18-mm and > 18 -mm branches of *C.*
375 *korshinskii*, respectively, and 1.1 mm, 1.6 mm, 2.0 mm and 2.4 mm for the branches of *S.*
376 *psammophila*, respectively.



377 The SF% of the two shrub species also increased with precipitation, but was inversely
378 proportional and gradually approached asymptotic values of 9.1% and 7.7% for *C.*
379 *korshinskii* and *S. psammophila*, respectively. As shown in Fig. 4, fast growth was evident
380 during rains ≤ 10 mm, but SF% slightly increased afterwards for both shrub species.

381

382 Fig. 4. Relationships of branch stemflow production (SF_b), shrub stemflow depth (SF_d) and
383 stemflow percentage (SF%) with precipitation amount (P) for *C. korshinskii* and *S.*
384 *psammophila*.

385

386 Precipitation amount was the most important factor affecting SFP and FR for *C.*
387 *korshinskii* and *S. psammophila*, but the most important biotic factor was different. BA was
388 the most influential plant trait that affected FR, and ILAB was the most important plant trait
389 affecting SFP during rains ≤ 10 mm. However, during heavy rain, BD and PBMS were the
390 most significant biotic factors for *C. korshinskii* and *S. psammophila*, respectively.

391

392 4 Discussion

393 4.1 Effective utilization of precipitation via stemflow production

394 Stemflow in *C. korshinskii* and *S. psammophila* increased with increasing precipitation
395 and branch size at both the branch (SF_b) and shrub scales (SF_d and SF%). However, *C.*
396 *korshinskii* had larger SF_b , SF_d and SF% values than did *S. psammophila* for all precipitation
397 categories. Although the greatest stemflow production was observed during rains > 20 mm for
398 the two shrub species, the inter-specific differences of SF_b , SF_d and SF% were highest at 3.2-,
399 8.5- and 8.3-fold larger for *C. korshinskii* during rains ≤ 2 mm, which indicated that *C.*
400 *korshinskii* utilized precipitation far more effectively during rains ≤ 2 mm at the branch and
401 shrub scale. These data indicate that stemflow was highly important for the survival of the
402 xerophytic shrubs in extreme drought. Additionally, *C. korshinskii* had a 2.8-fold larger SF_b
403 than that of *S. psammophila* for the 5–10-mm branches. Therefore, compared with *S.*



404 *psammophila*, more effectively might *C. korshinskii* utilize precipitation via greater stemflow
405 production, particularly the 5–10-mm young shoots during rains ≤ 2 mm.

406 The FR values indicated the efficiency with which individual branches could intercept
407 and channel raindrops (Siegert and Levia, 2014), thus leading to greater stemflow production.
408 The average FR of *S. psammophila* was 69.3, which agreed well with the 69.4 of *S.*
409 *psammophila* in the Mu Us sandland in China (Yang et al., 2008). The average FR for *C.*
410 *korshinskii* was 173.3, in contrast to the values of 156.1 (Jian et al., 2014) and 153.5 (Li et al.,
411 2008) for *C. korshinskii* in the western Loess Plateau of China. Furthermore, these two shrub
412 species had a larger FR than those of many other endemic xerophytic shrubs from
413 water-stressed ecosystems, e.g., *Tamarix ramosissima* (24.8) (Li et al., 2008), *Artemisia*
414 *sphaerocephala* (41.5) (Yang et al., 2008), *Reaumuria soongorica* (53.2) (Li et al., 2008),
415 *Hippophae rhamnoides* (62.2) (Jian et al., 2014). Therefore, both *C. korshinskii* and *S.*
416 *psammophila* utilized precipitation in a relatively efficient manner by producing stemflow,
417 and *C. korshinskii* produced stemflow more efficiently. The FR-specific difference achieved a
418 maximum of 3.5-fold larger for *C. korshinskii* during rains ≤ 2 mm and decreased with
419 increasing precipitation to 2.0-fold larger during rains > 20 mm.

420 SFP characterized stemflow production in terms of energy-conservation. *C. korshinskii*
421 had a larger SFP than *S. psammophila* for all the precipitation and BD categories, and during
422 rains ≤ 2 mm, the SFP-specific difference was maximized to 2.5-fold larger for *C. korshinskii*.
423 Additionally, the 5–10-mm branches had the largest average SFP of 2.2 mL g⁻¹ and 1.6
424 mL g⁻¹ in return, which, during rains > 20 mm, was maximized to 5.6 mL g⁻¹ and 4.6 mL g⁻¹
425 for *C. korshinskii* and *S. psammophila*, respectively (Table 3). Investing biomass into young
426 shoots provides considerable water benefits for xerophytic shrubs. Therefore, compared with
427 *S. psammophila*, more efficiently might *C. korshinskii* utilize precipitation by producing
428 greater stemflow, particularly for 5–10-mm young shoots during rains ≤ 2 mm.



429 Stemflow may preferentially incorporate precipitation into the rhizosphere, retaining it as
430 relatively stable soil moisture (Martinez-Meza and Whitford, 1996) and increasing drought
431 tolerance, particularly during long periods without rain. It was particularly significant that
432 young shoots were favoured in the presence of a greater water supply. Greater stemflow
433 production provided *C. korshinskii* with greater drought tolerance and a competitive edge in
434 water-stressed ecosystems.

435

436 **4.2 Utilization of more rains via a low precipitation threshold to start stemflow**

437 Precipitation below the threshold wet the canopy and then evaporated, so it did not
438 generate stemflow. The ≤ 2.5 -mm rains were entirely intercepted and evaporated to the
439 atmosphere for the xerophytic Ashe juniper communities at the central Texas of USA (Owens
440 et al., 2006), as well as most of the ≤ 5 -mm rains, particularly at the beginning raining stage
441 for xerophytic shrubs (*S. psammophila*, *Hedysarum scoparium*, *A. sphaerocephala* and
442 *Artemisia ordosica*) at the Mu Us sandland of China (Yang, 2010). The precipitation
443 threshold varied with factors such as the eco-zone, the PFT, the canopy structure, and the
444 branch architecture. A greater precipitation threshold partly explained why the SF% of trees
445 was smaller than that of shrubs (Llorens and Domingo, 2007). Particularly, the precipitation
446 threshold of xerophytic shrub species was as small as 0.3 mm for *T. vulgaris* at the northern
447 Lomo Herrero of Spain (Belmonte and Romero, 1998), but up to 2.7 mm for *A. farnesiana* at
448 Linares of Mexico (Návar and Bryan, 1990). In this study, at least a 0.9-mm rainfall was
449 necessary to initiate stemflow in *C. korshinskii*, which was in the range of 0.4-1.4 mm at the
450 precipitation threshold for *C. korshinskii* (Li et al., 2009; Wang et al., 2014). This result was
451 consistent with the 0.8 mm for *R. officinalis* at the northern Lomo Herrero of Spain (Belmont
452 and Romero, 1998) and 0.6 mm for *M. squamosa* at Qinghai-Tibet plateau of China (Zhang et
453 al., 2015). Comparatively, *S. psammophila* needed a 2.1-mm precipitation threshold to initiate



454 stemflow, which was consistent with the 2.2 mm threshold of *S. psammophila* in the Mu Us
455 desert (Li et al., 2009) and the 1.9 mm threshold for *R. soongorica* at the west of Loess
456 Plateau (Li et al., 2008) and the 1.8 mm threshold for *A. ordosica* at the Tengger desert of
457 China (Wang et al., 2014). Generally, for many xerophytic shrub species, the precipitation
458 threshold usually ranges between 0.4-2.2 mm, which is in accordance with the findings for
459 stemflow production (SF_b , SF_d and $SF\%$) and the production efficiency (SFP and FR), thus
460 indicating that rains ≤ 2 mm were particularly significant for the endemic plants in
461 water-stressed ecosystems.

462 Scant rainfall was the most prevalent type in arid and semiarid regions. Rains ≤ 5 mm
463 accounted for 74.8% of the annual rainfall events and 27.7% of the annual precipitation
464 amount at the Anjiapo catchment in the western Loess Plateau of China (with a MAP of 420
465 mm) (Jian et al., 2014). While at Haizetan in the south of Mu Us sandland of China (with a
466 MAP of 394.7 mm), rains ≤ 5 mm accounted for 49.0% of all the rainfall events and 13.8% of
467 the total precipitation amount of rainy season (lasting from May to September) (Yang 2010).
468 Additionally, rains ≤ 2.54 mm accounted for 60% of the total rainfall events and 5.4% of the
469 total precipitation amount at the eastern Edwards Plateau, the central Texas of USA (with a
470 MAP of 600-900 mm) (Owens et al., 2006). In this study, rains ≤ 2 mm accounted for 45.7%
471 of all the rainfall events and 7.2% of the precipitation amount during the 2014 and 2015 rainy
472 seasons. In general, *C. korshinskii* and *S. psammophila* produced stemflow during 71 (75.5%
473 of the total rainfall events) and 51 rainfall events (54.3% of the total rainfall events),
474 respectively. Because the precipitation threshold for *S. psammophila* was 2.1 mm, 20 rainfall
475 events of 12-mm, which encompassed 21.3% of all rainfall events, did not produce stemflow,
476 but stemflow production under these water stress conditions was an extra benefit for *C.*
477 *korshinskii*. Although the total amount was limited, it was of significant importance for the
478 survival of the xerophytic shrubs, particularly during long intervals with no rainfall.



479 In addition to the meteorological characteristics, the canopy structure and branch
480 architecture partly explained the inter-specific differences in the precipitation threshold
481 (Crockford and Richardson, 2000; Levia and Frost, 2003). A large, tall canopy created a large
482 rainfall interception area, also known as “canopy exposure” (Iida et al. 2011), particularly
483 during windy conditions (Van Stan et al, 2011). However, this advantage in stemflow
484 production might be offset by more consumption for wetting canopy and evaporation before
485 stemflow is generated in arid and semiarid regions, in which considerable evapotranspiration
486 potentially occurs. This phenomenon might be responsible for the smaller precipitation
487 threshold for stemflow production in *C. korshinskii*, which had a canopy height of 2.1 ± 0.2
488 m and a canopy area of $5.14 \pm 0.26 \text{ m}^2$, than *S. psammophila*, which had a canopy height of
489 3.5 ± 0.2 m and a canopy area of $21.35 \pm 5.21 \text{ m}^2$. Additionally, the canopy structure and
490 branch architecture also affected the water holding capacity (Herwitz, 1985), the interception
491 loss (Dunkerley, 2000), and consequently the precipitation threshold for stemflow generation
492 (Staelens et al., 2008). Nevertheless, the most influential plant traits had not determined yet,
493 and further stemflow studies was required at the finer leaf scale and temporal scale in the
494 future (Levia and Germer, 2015).

495

496 **4.3 Secure stemflow production advantage via beneficial leaf traits**

497 Further studies at the leaf scale indicated that leaf traits had a significant influence on
498 stemflow (Návar and Bryan, 1990; Carlyle-Mose, 2004; Garcia-Estringana et al., 2010). At
499 the individual shrub scale, the canopy gap, as represented by the LAI and the leaf mass,
500 provided direct access for raindrops to the branch surface (Crockford and Richardson, 2000).
501 The positive effects of LAI (Liang et al., 2009) and leaf biomass (Yuan et al., 2016) have
502 already been confirmed for *Stewartia monadelphpha* and *S. psammophila*, respectively. In a
503 study of European beech saplings, Levia et al. (2015) assumed that a threshold number of



504 leaves might exist for stemflow production. The positive effects could become negative if too
505 many leaves enclose the branches, which would benefit throughfall instead. In general,
506 factors such as a relatively large number of leaves (Li and Xiao, 2016), a large leaf area (Li et
507 al., 2015), a scale-like leaf arrangement (Owens et al., 2006), a small individual leaf area
508 (Sellin et al., 2012), a concave leaf shape (Xu et al., 2005), a densely veined leaf structure,
509 an upward leaf orientation (Crockford and Richardson, 2000), leaf pubescence
510 (Garcia-Estringana et al., 2010), and the leaf epidermis microrelief (e.g., the non-hydrophobic
511 leaf surface and the grooves within it) (Roth-Nebelsick et al., 2012) together result in the
512 retention of a large amount of precipitation in the canopy, supplying water for stemflow
513 production, and providing a beneficial morphology that enables the leaves to function as a
514 highly efficient natural water collecting and channelling system.

515 According to the field observations in this study, *C. korshinskii* had better leaf
516 morphology for stemflow production than did *S. psammophila*, owing to a lanceolate and
517 concave leaf shape, a pinnate compound leaf arrangement and a densely sericeous pressed
518 pubescence (Fig. 2). Additionally, experimental measurements indicated that *C. korshinskii*
519 had a larger MTA, LAB, LNB and SLW (an average of 54.4°, 2509.05 cm², 12479 and 126.04
520 g cm⁻², respectively) and a smaller ILAB (an average of 21.94 mm²) than did *S. psammophila*
521 (an average of 48.5°, 1797.93 cm², 2404, 73.87 g cm⁻² and 87.52 mm², respectively). The
522 larger SLW indicated that more biomass was deposited per unit leaf area. The concave leaf
523 shape, upward leaf orientation (MTA) and densely veined leaf structure (ILAB) (Xu et al.,
524 2005) provided stronger leaf structural support in *C. korshinskii* for the interception and
525 transportation of precipitation, particularly during highly intense rains. Therefore, in addition
526 to the leaf morphology, *C. korshinskii* was also equipped with more beneficial leaf structural
527 characteristics for stemflow production.

528 However, given that BML had strong effects on stemflow in *S. psammophila* (Yuan et al.,



2016), why were stem traits identified as the single most influential traits for stemflow
production in *C. korshinskii*, as indicated by the BMS in this study? The answer may partly
lie in the values of HV and PBMS. HV was computed as the cross-sectional area of the xylem
divided by the total leaf area supported by the stems (Sellin et al., 2012). A higher HV
indicates a potentially better water supply to leaves in terms of hydraulic conductance.
However, it could also be interpreted as indicating that more stem tissues are required to
support the unit leaf area for the normal function of the individual branch. The average HV of
C. korshinskii was 0.0507 and increased from 0.0438 for the 5–10-mm branches to 0.0615 for
the >18-mm branches and was an order of magnitude higher than in *S. psammophila*, which
averaged 0.0009 and remained nearly the same for different BD categories. The optimal
partitioning theory indicates that plants preferentially allocate biomass into the organs that
harvest the most limiting resource (Thornley, 1972; Bloom et al., 1985) and finally reach the
“functional equilibrium” of biomass allocation (Brouwer, 1963; Iwasa and Roughgarden,
1984). Therefore, a greater stem biomass might be required by *C. korshinskii* to support leaf
development than in *S. psammophila*, thus allowing more carbohydrate produced and
raindrops intercepted at the canopy. This possibility is consistent with the biomass allocation
patterns and leaf areas of the shrub species in this study. *C. korshinskii* allocated more
biomass into the stems with an average of PBMS of 85.6% and had a larger leaf area with an
average of LAB of 2509.1 cm² than *S. psammophila*, which had an average PBMS and LAB
of 81.9% and 1797.9 cm², respectively. The larger values of PBMS and LAB in *C.*
korshinskii were observed for all BD categories (Table 1). Additionally, the larger PBMS
helped to prevent the intercepted rain drops from falling off under windy conditions, which
also benefited stemflow production in *C. korshinskii*.

552

553 **5 Conclusions**



554 Compared with *S. psammophila*, *C. korshinskii* produced a larger amount of stemflow
555 more efficiently; an average 1.9, 1.3, 1.4, 1.6 and 2.5-fold increase in *C. korshinskii* was
556 observed for the branch stemflow production (SF_b), the shrub stemflow depth (SF_d), the shrub
557 stemflow percentage (SF%), the stemflow productivity (SFP) and the stemflow funnelling
558 ratio (FR), respectively. The largest specific difference in stemflow production (SF_b , SF_d and
559 SF%) and the production efficiency (SFP and FR) was during rains ≤ 2 mm, which were the
560 most frequent rainfall events. Although the total amount of rainfall was limited, it was of
561 great importance for *C. korshinskii* to survive and thrive, particularly during the extreme
562 drought period. Additionally, the inter-specific differences in SF_b , SF_d , SF% and SFP were
563 maximized for the 5–10-mm branches; this result was particularly significant because it
564 encouraged young shoots by supplying more water.

565 Beneficial leaf traits, including a lanceolate and concaved leaf shape, a pinnate
566 compound leaf arrangement, a densely sericeous pressed pubescence, an upward leaf
567 orientation (MTA), a large leaf area (LAB), a relatively large number of leaves (LNB), a large
568 leaf area index (LAI), a small individual leaf area (ILAB), and a large specific leaf weight
569 (SLW), might be responsible for the superior stemflow production in *C. korshinskii*. Along
570 with the canopy structure, these leaf traits may account for the lower precipitation threshold
571 to initiate stemflow in *C. korshinskii* (0.9 mm) than in *S. psammophila* (2.1 mm). A lower
572 precipitation threshold enabled *C. korshinskii* to harvest more water from rainfall via
573 stemflow.

574 In conclusion, a higher and more efficient stemflow, a lower precipitation threshold and
575 beneficial leaf traits provided *C. korshinskii* with greater drought tolerance and a competitive
576 edge in a water-stressed ecosystem.

577

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584

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793 2008.



794 **Table captions**

795

796 **Table 1.** Comparison of leaf traits, branch morphology and biomass indicators of *C.*
797 *korshinskii* and *S. psammophila*.

798

799 **Table 2.** Comparison of stemflow production (SF_b , SF_d and $SF\%$) between *C. korshinskii* and
800 *S. psammophila*.

801

802 **Table 3.** Comparison of stemflow productivity (SFP) between *C. korshinskii* and *S.*
803 *psammophila*.

804

805 **Table 4.** Comparison of the funneling ratio (FR) for *C. korshinskii* and *S. psammophila*.



806

Table 1. Comparison of leaf traits, branch morphology and biomass indicators of *C. korshinskii* and *S. psammophila*.

| Plant traits | <i>C. korshinskii</i> (categorized by BD, mm) | | | | | <i>S. psammophila</i> (categorized by BD, mm) | | | | | |
|--------------------|---|--------|--------|--------|-----------|---|--------|--------|--------|-----------|-------------------|
| | 5-10 | 10-15 | 15-18 | >18 | Avg. (BD) | 5-10 | 10-15 | 15-18 | >18 | Avg. (BD) | |
| Leaf traits | LAB (cm ²) | 1202.7 | 2394.5 | 3791.2 | 5195.2 | 2509.1 ±1355.3 | 499.2 | 1317.7 | 2515.2 | 3533.6 | 1797.9 ±1118.0 |
| | LNB | 4787 | 11326 | 20071 | 29802 | 12479 ±8409 | 392 | 1456 | 3478 | 5551 | 2404 ±1922 |
| | ILAB (mm ²) | 25.4 | 21.3 | 18.9 | 17.5 | 21.9 ±3.0 | 135.1 | 93.1 | 72.6 | 64.3 | 93.1 ±27.8 |
| | SLW (g cm ⁻²) | 126.4 | 126.0 | 125.7 | 125.6 | 126.0 ±0.3 | 95.6 | 74.5 | 63.0 | 58.1 | 73.9 ±14.5 |
| | HV | 0.0438 | 0.0513 | 0.0572 | 0.0615 | 0.0507 ±0.0064 | 0.0010 | 0.0009 | 0.0009 | 0.0009 | 0.0009 ±0.0001 |
| Branch morphology | BD (mm) | 8.17 | 12.49 | 16.61 | 20.16 | 12.48 ±4.16 | 7.91 | 12.48 | 16.92 | 19.76 | 13.73 ±4.36 |
| | BL (cm) | 137.9 | 160.3 | 195.9 | 200.7 | 161.5 ±35.0 | 212.5 | 260.2 | 290.4 | 320.1 | 267.3 ±49.7 |
| | BA (°) | 63 | 56 | 63 | 64 | 60 ±18 | 64 | 63 | 51 | 60 | 60 ±20 |
| Biomass indicators | BML (g) | 13.9 | 19.0 | 30.2 | 41.4 | 19.9 ±10.8 | 5.4 | 18.0 | 40.0 | 61.3 | 27.9 ±20.7 |
| | BMS (g) | 62.9 | 121.4 | 236.4 | 375.8 | 141.1 ±110.8 | 23.0 | 81.4 | 188.5 | 295.5 | 130.7 ±101.4 |
| | PBMS (%) | 82.0 | 86.3 | 88.7 | 90.0 | 85.6 ±3.1 | 80.8 | 81.8 | 82.5 | 82.8 | 81.9 ±0.8 |

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Note: LAB and LNB are leaf area and number of branch, respectively. ILAB is individual leaf area of branch. SLW is the specific leaf weight, and HV was the Huber value. BD, BL and BA are average branch basal diameter, length and angle, respectively. BML and BMS are biomass of leaves and stems, respectively. PBMS is the percentage of leaf biomass to that of branch. The average values mentioned above are expressed as the means ±SE.



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Table 2. Comparison of stemflow production (SF_b , SF_d and $SF\%$) between *C. korshinskii* and *S. psammophila*.

| Intra- and inter-specific differences | Stemflow indicators | BD categories (mm) | Precipitation categories (mm) | | | | | | Avg.(P) |
|---|---------------------|--------------------|-------------------------------|-------|-------|-------|--------|--------|---------|
| | | | ≤2 | 2-5 | 5-10 | 10-15 | 15-20 | >20 | |
| Intra-specific differences in <i>C. korshinskii</i> (CK) | SF_b (mL) | 5-10 | 10.7 | 29.8 | 73.5 | 109.9 | 227.6 | 306.1 | 119.0 |
| | | 10-15 | 26.0 | 64.0 | 166.1 | 236.0 | 478.6 | 689.7 | 262.4 |
| | | 15-18 | 44.3 | 103.3 | 279.9 | 416.6 | 826.0 | 1272.3 | 464.5 |
| | | >18 | 69.5 | 145.4 | 424.4 | 631.4 | 1226.9 | 1811.7 | 679.9 |
| | | Avg.(BD) | 28.4 | 67.3 | 180.6 | 264.6 | 529.2 | 771.4 | 290.6 |
| | SF_d (mm) | N/A | 0.09 | 0.24 | 0.63 | 0.91 | 1.85 | 2.64 | 1.00 |
| | $SF\%$ (%) | N/A | 5.8 | 6.6 | 8.8 | 7.5 | 10.1 | 8.9 | 8.0 |
| Intra-specific differences in <i>S. psammophila</i> (SP) | SF_b (mL) | 5-10 | 2.8 | 8.9 | 28.8 | 47.2 | 66.5 | 120.0 | 43.0 |
| | | 10-15 | 7.6 | 23.2 | 76.6 | 134.6 | 188.3 | 353.5 | 121.8 |
| | | 15-18 | 12.0 | 35.9 | 121.6 | 223.4 | 319.4 | 592.6 | 201.5 |
| | | >18 | 16.2 | 52.3 | 165.5 | 289.2 | 439.6 | 860.4 | 281.8 |
| | | Avg.(BD) | 9.0 | 28.0 | 91.6 | 162.2 | 234.8 | 444.3 | 150.3 |
| | SF_d (mm) | N/A | 0.01 | 0.11 | 0.48 | 0.89 | 1.27 | 2.23 | 0.78 |
| | $SF\%$ (%) | N/A | 0.7 | 3.0 | 6.1 | 6.8 | 7.2 | 7.9 | 5.5 |
| Inter-specific differences (the ratio of the stemflow production of CK to that of SP) | SF_b | 5-10 | 3.8 | 3.3 | 2.6 | 2.3 | 3.4 | 2.6 | 2.8 |
| | | 10-15 | 3.4 | 2.8 | 2.2 | 1.8 | 2.5 | 2.0 | 2.2 |
| | | 15-18 | 3.7 | 2.9 | 2.3 | 1.9 | 2.6 | 2.2 | 2.3 |
| | | >18 | 4.3 | 2.8 | 2.6 | 2.2 | 2.8 | 2.1 | 2.4 |
| | Avg.(BD) | 3.2 | 2.4 | 2.0 | 1.6 | 2.3 | 1.7 | 1.9 | |
| | SF_d | N/A | 8.5 | 2.2 | 1.3 | 1.0 | 1.5 | 1.2 | 1.3 |
| | $SF\%$ | N/A | 8.3 | 2.2 | 1.4 | 1.1 | 1.4 | 1.1 | 1.4 |

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Note: BD is the branch basal diameter; P is the precipitation amount; CK and SP are the abbreviations of *C. korshinskii* and *S. psammophila*, respectively.



814

Table 3. Comparison of stemflow productivity (SFP) between *C. korshinskii* and *S. psammophila*.

| Intra- and inter-specific differences | BD categories (mm) | Precipitation categories (mm) | | | | | | Avg.(P) |
|--|--------------------|-------------------------------|------|------|-------|-------|------|---------|
| | | ≤2 | 2-5 | 5-10 | 10-15 | 15-20 | >20 | |
| Intra-specific differences in <i>C. korshinskii</i> (CK) (mL g ⁻¹) | 5-10 | 0.20 | 0.56 | 1.37 | 2.04 | 4.18 | 5.60 | 2.19 |
| | 10-15 | 0.19 | 0.47 | 1.20 | 1.72 | 3.47 | 4.96 | 1.90 |
| | 15-18 | 0.17 | 0.38 | 1.05 | 1.55 | 3.08 | 4.74 | 1.73 |
| | >18 | 0.15 | 0.35 | 1.00 | 1.46 | 2.95 | 4.35 | 1.62 |
| | Avg.(BD) | 0.19 | 0.47 | 1.21 | 1.78 | 3.60 | 5.08 | 1.95 |
| Intra-specific differences in <i>S. psammophila</i> (SP) (mL g ⁻¹) | 5-10 | 0.11 | 0.34 | 1.10 | 1.83 | 2.51 | 4.59 | 1.64 |
| | 10-15 | 0.08 | 0.25 | 0.82 | 1.43 | 1.98 | 3.72 | 1.29 |
| | 15-18 | 0.05 | 0.16 | 0.53 | 0.97 | 1.40 | 2.61 | 0.88 |
| | >18 | 0.05 | 0.15 | 0.47 | 0.82 | 1.25 | 2.44 | 0.80 |
| | Avg.(BD) | 0.07 | 0.23 | 0.76 | 1.31 | 1.84 | 3.43 | 1.19 |
| Inter-specific differences (the ratio of the SFP values of CK to that of SP) | 5-10 | 1.8 | 1.7 | 1.3 | 1.1 | 1.7 | 1.2 | 1.3 |
| | 10-15 | 2.4 | 1.9 | 1.5 | 1.2 | 1.8 | 1.3 | 1.5 |
| | 15-18 | 2.8 | 2.4 | 2.0 | 1.6 | 2.2 | 1.8 | 2.0 |
| | >18 | 3.0 | 2.3 | 2.1 | 1.8 | 2.4 | 1.8 | 2.0 |
| | Avg.(BD) | 2.7 | 2.0 | 1.6 | 1.4 | 2.0 | 1.5 | 1.6 |

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Note: BD is the branch basal diameter; P is the precipitation amount; CK and SP are the abbreviations of *C. korshinskii* and *S. psammophila*, respectively.



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Table 4. Comparison of the funneling ratio (FR) for *C. korshinskii* and *S. psammophila*.

| Intra- and inter-specific differences | BA categories (°) | Precipitation categories (mm) | | | | | | Avg.(P) |
|---|-------------------|-------------------------------|--------|--------|--------|--------|--------|---------|
| | | ≤2 | 2-5 | 5-10 | 10-15 | 15-20 | >20 | |
| Intra-specific differences in <i>C. korshinskii</i> (CK) | ≤30 | 100.18 | 127.68 | 168.14 | 125.30 | 193.06 | 170.31 | 149.90 |
| | 30-60 | 125.89 | 133.77 | 178.5 | 157.84 | 205.19 | 182.07 | 164.65 |
| | 60-80 | 135.51 | 148.94 | 192.45 | 165.83 | 217.03 | 188.64 | 176.06 |
| | >80 | 133.17 | 167.44 | 205.53 | 182.61 | 276.02 | 226.08 | 198.16 |
| | Avg.(BA) | 129.17 | 144.84 | 187.74 | 162.34 | 219.61 | 190.34 | 173.34 |
| Intra-specific differences in <i>S. psammophila</i> (SP) | ≤30 | 32.60 | 37.33 | 52.02 | 59.00 | 65.75 | 85.19 | 54.97 |
| | 30-60 | 34.50 | 43.44 | 65.67 | 70.63 | 77.74 | 92.28 | 64.78 |
| | 60-80 | 37.83 | 47.92 | 77.99 | 78.41 | 82.31 | 97.72 | 72.39 |
| | >80 | 44.88 | 54.99 | 93.45 | 94.74 | 94.09 | 115.72 | 85.57 |
| | Avg.(BA) | 36.65 | 46.01 | 72.57 | 75.34 | 80.45 | 96.09 | 69.25 |
| Inter-specific differences (the ratio of the FR values of CK to that of SP) | ≤30 | 3.1 | 3.4 | 3.2 | 2.1 | 2.9 | 2.0 | 2.7 |
| | 30-60 | 3.7 | 3.1 | 2.7 | 2.2 | 2.6 | 2.0 | 2.5 |
| | 60-80 | 3.6 | 3.1 | 2.5 | 2.1 | 2.6 | 1.9 | 2.4 |
| | >80 | 3.0 | 3.0 | 2.2 | 1.9 | 2.9 | 2.0 | 2.3 |
| | Avg.(BA) | 3.5 | 3.2 | 2.6 | 2.2 | 2.7 | 2.0 | 2.5 |

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Note: BA is the branch inclined angle; P is the precipitation amount; CK and SP are the abbreviations of *C. korshinskii* and *S. psammophila*, respectively.



820 **Figure captions**

821

822 **Fig. 1.** Location of the experimental stands and facilities for stemflow measurements of *C.*
823 *korshinskii* and *S. psammophila* at the Liudaogou catchment in the Loess Plateau of
824 China.

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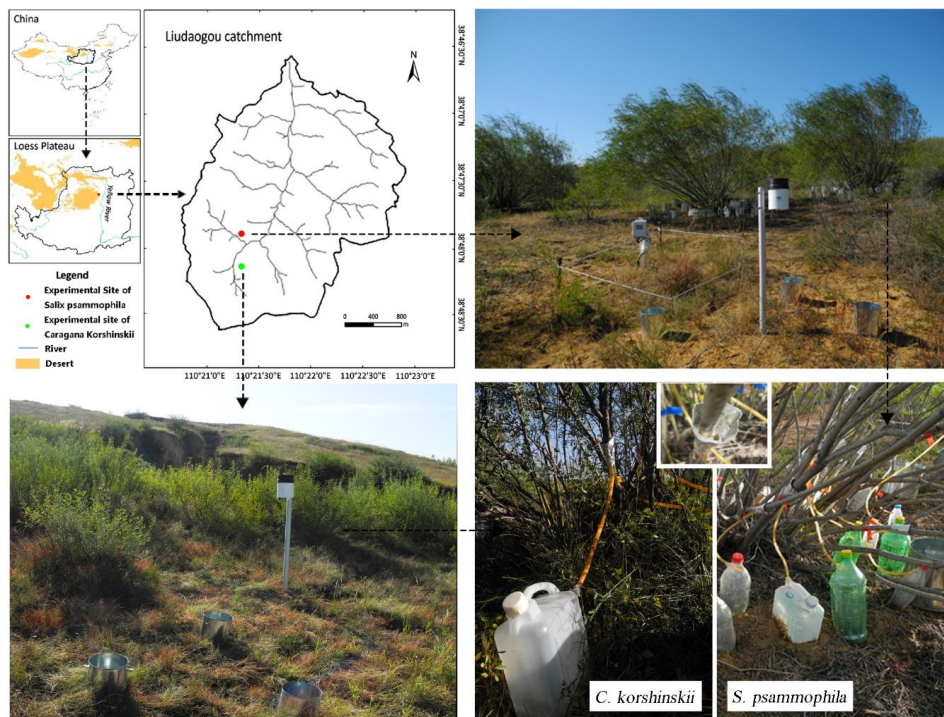
826 **Fig. 2.** Comparison of leaf morphologies of *C. korshinskii* and *S. psammophila*.

827

828 **Fig. 3.** Verification of the allometric models for estimating the biomass and leaf traits of *C.*
829 *korshinskii*. BML and BMS refer to the biomass of the leaves and stems, respectively,
830 and LAB and LNB refer to the leaf area and the number of branches, respectively.

831

832 **Fig. 4.** Relationships of branch stemflow production (SF_b), shrub stemflow depth (SF_d) and
833 stemflow percentage ($SF\%$) with precipitation amount (P) for *C. korshinskii* and *S.*
834 *psammophila*.



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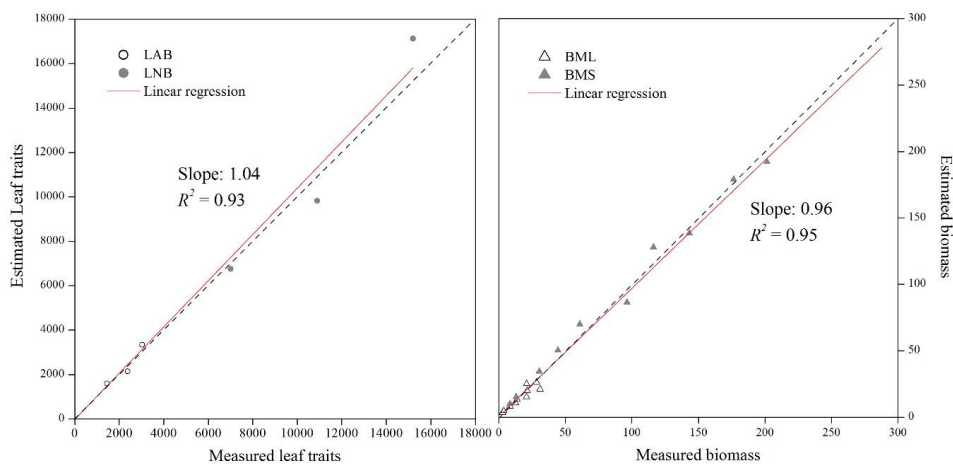
Fig. 1. Location of the experimental stands and facilities for stemflow measurements of *C. korshinskii* and *S. psammophila* at the Liudaogou catchment in the Loess Plateau of China.



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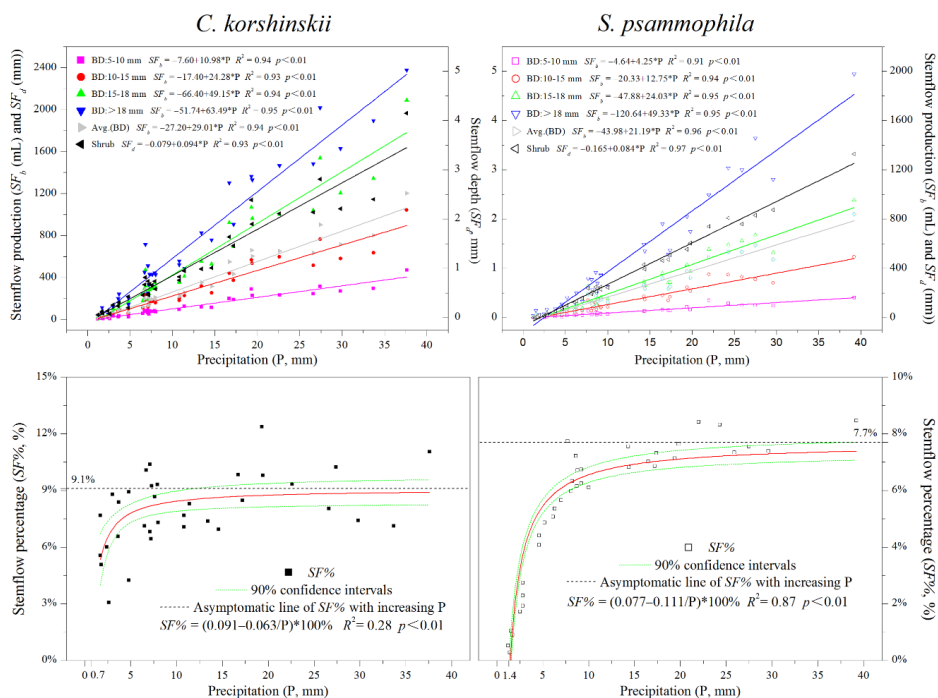
839

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840

841 **Fig. 3.** Verification of the allometric models for estimating the biomass and leaf traits of *C.*
842 *korshinskii*. BML and BMS refer to the biomass of the leaves and stems, respectively, and
843 LAB and LNB refer to the leaf area and the number of branches, respectively.



844

845 **Fig. 4.** Relationships of branch stemflow production (SF_b), shrub stemflow depth (SF_d) and
 846 stemflow percentage ($SF\%$) with precipitation amount (P) for *C. korshinskii* and
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