

1 **Effects of Superhydrophobic Sand Mulching on**
2 **Evapotranspiration and Phenotypic Responses in Tomatoes**
3 **(*Solanum lycopersicum*) under Normal and Reduced Irrigation**

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12
13 **Abstract**

14 Irrigated agriculture in arid and semi-arid regions is a vital contributor to the global food
15 supply; however, these regions endure massive evaporative losses that are compensated by
16 unsustainable freshwater withdrawals. Plastic mulches have been used to curtail
17 evaporation, improve water-use efficiency, and ensure food–water security, but they are
18 non-biodegradable and their disposal is unsustainable. We recently developed
19 superhydrophobic sand (SHS), which comprises sand grains with a nanoscale wax coating
20 that could offer a more sustainable mulching solution. Here, the effects of adding a 1.0 cm-
21 thick layer of SHS mulch on the evapotranspiration and phenotypic responses of tomato
22 (*Solanum lycopersicum*) plants are studied under normal and reduced irrigation. Under
23 both irrigation regimes, SHS mulching suppressed evaporation and enhanced transpiration
24 by 78% and 17%, respectively relative to the bare soil. Overall, SHS mulching enhanced
25 root xylem vessel diameter, stomatal aperture, stomatal conductance, and chlorophyll
26 content index by 21%, 25%, 28%, and 23%, respectively. Total fruit yields, total dry mass,
27 and harvest index increased in SHS-mulched plants by 33%, 20%, and 16%, respectively
28 than in bare soil. These findings demonstrate the potential of SHS to boost irrigation
29 efficiency in water-limited environments and provide mechanistic insights behind yield
30 enhancement by SHS mulching.

31

32 **KEYWORDS:** evapotranspiration; *Solanum lycopersicum*; superhydrophobic sand;
33 transpiration efficiency; chlorophyll content index; stomatal conductance; harvest index.

34

35

36 **INTRODUCTION**

37 Irrigated agricultural lands are of vital importance, as they represent 20% of the total
38 cultivated land but contribute over 40% of the global food production (WWAP, 2009).

39 Irrigated agriculture has been expanding to feed the growing global population; for
40 example, India, China, and Brazil have witnessed a 30% (Jain et al., 2020), 52% (Zhu et
41 al., 2013), and 52% (Carvalho et al., 2020) increase in cultivated areas under irrigation in

42 the last 50 years, respectively. Further, between 2012 and 2030, the global irrigated land
43 area is expected to increase by over $\approx 30\%$, from 310 Mha to 402 Mha (Darko et al., 2016).

44 These food production trends are sustainable only when freshwater is abundant. However,
45 groundwater resources have been extensively exploited for irrigation in densely populated
46 arid and semi-arid regions and are thus rapidly declining (Steward et al., 2013, Watto et
47 al., 2018b, Watto et al., 2018a, Scanlon et al., 2012, Famiglietti, 2014). Therefore, water
48 consumption patterns in irrigated agriculture must be studied, and new technologies must
49 be created to achieve global water–food security.

50 Water applied to topsoil is lost via evaporation, transpiration, and percolation
51 (Sutanto, 2012). High temperatures and dry winds in arid and semi-arid regions lead to
52 substantial evaporative and transpiration losses (Al-Naizy, 2012, Balugani et al., 2017),
53 whereas percolation is due to the poor water-holding capacity of sandy soils (Lehmann et
54 al., 2019, Or and Lehmann, 2019). These water losses are compensated via irrigation that,
55 due to the sheer size of agricultural operations, claims the lion’s share (approximately
56 $\approx 80\%$) of global annual freshwater consumption (2500 km³/yr) (Shiklomanov, 2000,
57 Hoekstra and Mekonnen, 2012). Whereas evaporation and percolation do not directly
58 contribute to photosynthesis during plant growth, transpiration is crucial in maintaining the
59 optimal temperature required for plants’ metabolic processes (Grill and Ziegler, 1998,
60 Hetherington and Woodward, 2003). Transpiration thus cannot be reduced while high crop
61 productivity is maintained without genetic engineering. Percolation could be reduced by
62 applying a water barrier layer beneath the plant root zone, but this process is labor-intensive

63 and expensive (Nkurunziza et al., 2019). Thus, many researchers have focused on reducing
64 evaporative losses.

65 Together, evaporation and transpiration constitute evapotranspiration (ET), a
66 process that plays an important role in irrigation and agriculture practices (Hatfield and
67 Dold, 2019, Rawitz and Hadas, 1994). The importance of transpiration is reflected in the
68 water-use efficiency (WUE) of plants, i.e., the ratio of the total biomass (root and shoot)
69 formed to the cumulative water transpired (Kadam et al., 2015), and the transpiration
70 efficiency (TE), i.e., the ratio of the shoot biomass produced to the cumulative water
71 transpired (Vadez et al., 2014). The TE represents an aspect of WUE that depends on the
72 water-conducting (i.e., hydraulic) potential of a plant to facilitate shoots' physiological
73 processes that enhance biomass under different soil water scenarios(Vadez et al., 2014).
74 Therefore, the TE is of interest in water-scarce environments where crops are most
75 sensitive to soil moisture (Condon et al., 2002, Sinclair, 2012). Transpiration also
76 influences the plants' reproductive efficiency, defined by the harvest index (HI) entailing
77 the ratio of total yield to total vegetative biomass produced, pinpointing the plant resources
78 invested in yield production (Porker et al., 2020, Unkovich et al., 2010). Hence, whereas
79 water uptake and efficient use of water are critical for plant growth and biomass
80 accumulation, the development and growth of reproductive organs determines the HI,
81 which is critical to crop yield (Hammer et al., 2021).

82 Notably, evaporation is a wasteful component of ET and should be minimized.
83 Mulching, i.e., applying a vapor diffusion barrier on the topsoil to curtail water
84 evaporation, has been proven to enhance soil moisture content and provide enhanced
85 transpiration (Moitra, 1996, Zhang et al., 2018, Farzi et al., 2017), plant biomass, and yields
86 (Ramalan, 2000, Zhang, 2017a, Mukherjee, 2010). Mulching has been demonstrated to
87 enhance leaf chlorophyll content (Wang et al., 2015), photosynthesis (Niu et al., 2020,
88 Zhang et al., 2019), and TE (Balwinder-Singh et al., 2011), as well as improve plant root
89 growth and roots' architectural and anatomical properties, such as root diameter and late
90 xylem vessel diameter, which improves water and nutrient uptake from the soil (Zhan et
91 al., 2019, Larsson and Jensen, 1996). Low-density polyethylene sheets approximately 0.1
92 mm thick have been used extensively for mulching in developed countries (Kasirajan and
93 Ngouajio, 2012). Their application in the developing world is also on the rise, e.g.,

94 consumption in China is set to exceed two million tons per year by 2024 (Liu et al., 2014,
95 Wang et al., 2013). Despite their benefits (Zhang, 2017b, Zhang et al., 2019, Balwinder-
96 Singh et al., 2011, Yin W, 2019, Li, 2004), plastics are non-biodegradable, and their
97 eventual disposal into landfills is unsustainable (Kasirajan and Ngouajio, 2012, Halley et
98 al., 2001). Recent findings on the leaching of phthalates from plastic mulches into soils and
99 their adverse effects on the soil quality and microbial activity over the long term are also
100 concerning (Shah and Wu, 2020). Therefore, sustainable mulching technologies are
101 needed.

102 We recently developed superhydrophobic sand (SHS) mulch technology (Mishra
103 et al., 2017, Gallo Jr et al., 2021a), a nature-inspired material comprising common sand
104 grains or sandy soils coated with a nanoscale layer of paraffin wax (wax to sand mass ratio
105 is 1:1000). The micro- and nanoscale surface roughness of the sand grains/particles and the
106 hydrophobic nature of wax induce superhydrophobicity (Arunachalam et al., 2019). When
107 laid on topsoil with a sub-surface irrigation system, a 5–10 mm-thick SHS layer can
108 insulate the wet soil from solar radiation and dry air, thereby acting as a diffusion barrier,
109 which reduces evaporation (Gallo Jr et al., 2021b). Multi-year field trials of SHS on
110 tomatoes (*Solanum lycopersicum*), wheat (*Triticum aestivum*), and barley (*Hordeum*
111 *vulgare*) under naturally arid land conditions in Saudi Arabia have revealed significant
112 enhancements in plant growth and yields (Gallo Jr et al., 2021a). Compared with plastic
113 mulch, SHS is cheaper and more environmentally friendly, as it is made from readily
114 abundant sands and paraffin wax that is easily degraded by soil microbes (Marino, 1998,
115 Roper, 2004). After nine months of use, the wax is decomposed by microbial activity and
116 solar radiation and incorporated into the soil after the crop cycle without affecting soil
117 microbial compositions, obviating landfilling (Gallo Jr et al., 2021a). Despite these
118 promising field results, quantitative insights into the effects of SHS on ET and the
119 phenotypic traits responsible for yield enhancement are lacking.

120 This work therefore aims to quantify the effects of mulching with SHS on tomato
121 plants by pinpointing the ET dynamics and consequent plant phenotypes, including plant
122 height, stomatal conductance, stomatal pore size (aperture), leaf chlorophyll content, fruit
123 yields, HI, TE, fresh mass, dry mass, xylem vessel and root diameter. These effects are

124 considered in plants grown in controlled growth chambers under normal (**N**) and reduced
125 (**R**) irrigation scenarios at 100% field capacity and 50% of field capacity, respectively.

126

127 **MATERIALS AND METHODS**

128 **Plants and SHS mulch**

129 The tomato plants (*Solanum lycopersicum*), Seminis variety (St. Louis, Missouri, US),
130 were purchased from local seed stores in Jeddah, Saudi Arabia. The SHS was produced
131 from common sand and paraffin wax following the optimized protocol detailed in previous
132 work (Mishra et al., 2017, Gallo Jr et al., 2021a); readers are referred to these works for
133 details on SHS production. Briefly, sand was added to an organic solvent containing
134 dissolved wax. The solvent was then removed by changing the pressure and temperature
135 and condensed for reuse, leaving behind the SHS.

136

137 **Plant growth conditions, treatment, and experimental design**

138 The tomato seeds were sown in plastic trays using a potting mix from Stender AG
139 (Schermbek, Germany) and grown in Percival growth chambers. After four weeks, the
140 seedlings were transplanted into pots 1,870 cm³ in volume (15 cm top diameter × 10.5 cm
141 bottom diameter × 14.5 cm height) containing approximately 2.4 kg of local sandy soil.
142 The pots were watered via sub-surface irrigation to two field capacity levels every two
143 days: 100% field capacity for **N** irrigation (i.e., the maximum soil moisture content after
144 drainage of excess water from fully saturated potted soil) and 50% of this field capacity for
145 **R** irrigation. The irrigation level of each pot was determined gravimetrically. Thirty-two
146 pots were prepared, including 16 with plants and 16 without plants, the latter of which were
147 used to quantify evaporative losses.

148 The pots were separated into two groups: those comprising SHS and those
149 containing only bare soil. In each group, half of the pots were subjected to **N** irrigation
150 while the other half was maintained under **R** irrigation. For pots containing SHS, a 1.0 cm-
151 thick layer of SHS was applied to the potting soil (approximately 265 g SHS/pot). Thus,
152 there were four treatment combinations (SHS-**N**, bare soil-**N**, SHS-**R**, and bare soil-**R**),
153 each with two replicates with and without plants. All pots were subjected to complete
154 randomization in a 2 × 2 factorial design involving two experimental factors (i.e., soil

155 mulching and irrigation regime), each with two levels and four replicates. During the
156 growth period, nutrient solutions were applied to each pot every two weeks comprising in
157 rates per kg of dry soil: 200 mg N/kg, 250 mg P/kg, 200 mg K/kg, 150 mg Ca/kg, 30 mg
158 S/kg, 2 mg Cu/kg, 4 mg Zn/kg, 3 mg Mn/kg, 0.5 mg B/kg, and 0.25 mg Mo/kg. The plants
159 were grown for 98 days under a 14/10 h light/dark photoperiod using artificial fluorescent
160 lighting at $800 \mu\text{mol m}^{-2} \text{s}^{-1}$ of photosynthetically active radiation, a day/night temperature
161 of $28/20^\circ\text{C} \pm 2^\circ\text{C}$, and $60 \pm 2\%$ air relative humidity.

162

163 **Evapotranspiration (ET)**

164 The ET was partitioned into evaporation and transpiration by performing gravimetric
165 measurements of pots every two days until the final harvest. The water loss from each pot
166 was monitored, and water was added to compensate for the water lost. The daily ET was
167 recorded as the total water lost from each pot containing plants between the time of
168 irrigation to the time of weighing as:

169

$$170 \quad ET = (\text{Initial weight of pot with plant} - \text{final weight of pot with plant}) / (\text{time between} \\ 171 \quad \text{measurements}). \quad (\text{Eq. i})$$

172

173 **Evaporation**

174 Under the assumption that the pots with and without plants experienced a similar amount
175 of evaporation, the daily water loss by evaporation was estimated using pots without plants
176 as:

177

$$178 \quad \text{Evaporation} = (\text{Initial weight of pot without plant} - \text{final weight of pot without plant}) / (\text{time} \\ 179 \quad \text{between measurements}). \quad (\text{Eq. ii})$$

180

181 **Transpiration**

182 The daily transpiration was determined as:

183

$$184 \quad \text{Transpiration} = (ET - \text{Evaporation}). \quad (\text{Eq. iii})$$

185

186 **Soil water content (SWC)**

187 The total available soil water content (SWC; i.e., the field capacity) was maintained at a
188 relatively constant level by compensating for the water lost through ET after each
189 gravimetric measurement scheduled at two-day interval, as has been done by prior
190 researchers (Ray and Sinclair, 1998, Pellegrino et al., 2004).

191

192 **Stomatal conductance and leaf chlorophyll content**

193 The leaf stomatal conductance was determined using an AP4 Porometer (Delta T,
194 Cambridge, UK). Measurements were performed on three young but fully expanded leaves
195 once a week (between 10:00 and 12:00), and the mean conductance for each treatment
196 combination was calculated. The leaf chlorophyll content index (CCI) was measured using
197 a CCM-200 Chlorophyll Content Meter (Optic-Sciences, Inc. Hudson NH03051, USA),
198 with measurements being performed on three young but fully expanded leaves.

199

200 **Leaf stomatal pore and root anatomical features**

201 Microscopic analyses of leaf stomata and root anatomic structures were performed on
202 samples collected on the final date of harvest. Two fully expanded upper leaves were cut
203 from each plant (between 11:00 and 11:30) and immediately immersed in 70% ethanol
204 inside 50 ml centrifuge tubes until they were analyzed. After two weeks, the preserved leaf
205 tissues were washed with deionized water three times, added to 40 ml of concentrated
206 sodium hypochlorite, and left to stay for approximately 4 hours. The leaf tissues were re-
207 washed with deionized water, and fresh 70% ethanol was again added onto the samples.
208 This clearing process removed the chlorophyll from the leaves and rendered them white in
209 appearance. The stomata of the cleared leaves were then observed using a digital
210 microscope (Leica DVM6) equipped with the tools to measure the length and width of the
211 stomatal pores. The total stomatal aperture area was calculated using the formula for the
212 area of an ellipse, as

213

214
$$\text{Stomatal aperture area} = \pi \times a \times b, \quad (\text{Eq. iv})$$

215

216 where a and b represent the semi-major axis (or radius) and semi-minor axis (or width) of
217 the ellipse, respectively.

218 Root samples 10 cm in length from the growing tip were cut using a razor, washed in
219 deionized water, and fixed in 70% ethanol until their microscopic analysis. Two hand-cut
220 root tissues were made per plant and their cross-sections were analyzed using the same
221 digital microscope to measure diameter of the xylem vessels and the total root.

222

223 **Plant growth, fruit yield, and biomass**

224 The plant height was measured every week during the 98-day experimental period.
225 Tomatoes were harvested and weighed at the time of harvest. During the final harvest, the
226 shoots and roots of each plant were weighed, and then put in paper bags and oven-dried at
227 105 °C for four days to determine their dry mass.

228

229 **Harvest index (HI) and transpiration efficiency (TE)**

230 After harvest, the HI was calculated for each plant as the ratio of total fruit yield (fruit mass
231 per plant) to total fresh mass (shoot and roots) per plant, as:

232

$$233 \quad HI = \frac{\text{Total fruit yield}}{\text{Total fresh biomass}}. \quad (\text{Eq. v})$$

234

235 The TE of each plant was calculated as the ratio of dry shoot biomass in grams to the
236 amount of water transpired in kilograms, as:

237

$$238 \quad TE = \frac{\text{Shoot dry biomass produced (g)}}{\text{mass of water transpired (kg)}}. \quad (\text{Eq. vi})$$

239

240 **Data analysis**

241 Data analysis and graphical presentations were performed using Origin Pro Software (2019
242 version). A three-way analysis of variance (ANOVA) was used to analyze the effects of
243 soil mulching, irrigation regimes, growth stage (i.e., time), and their interactions on ET.
244 For the variables averaged across time (i.e., time-independent), a two-way factorial
245 ANOVA was performed to determine the effects of soil mulching, irrigation regimes, and

246 their interactions. In each analysis, mean comparisons were done using the Tukey test at
247 the $p < 0.05$ level of statistical significance.

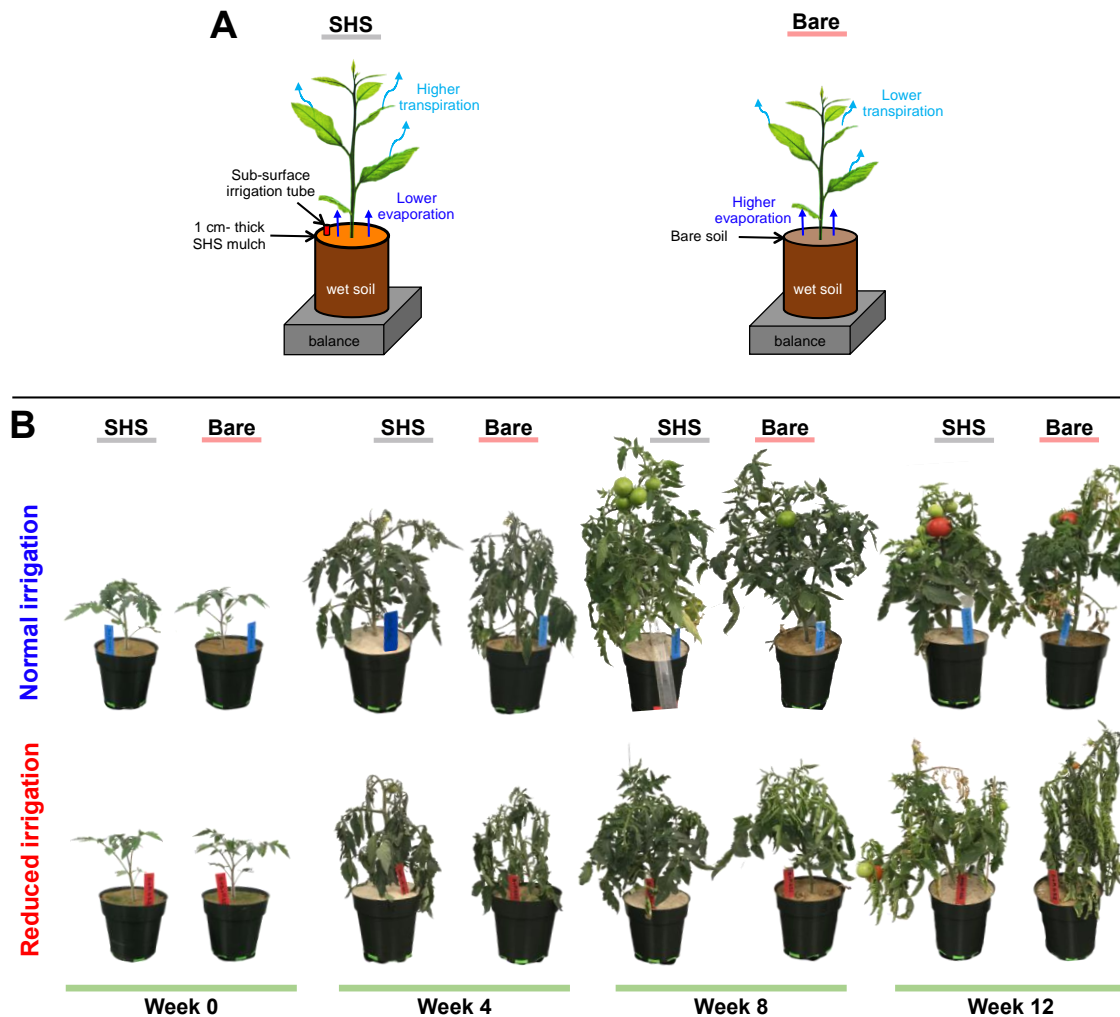
248

249 RESULTS

250 Plant growth

251 Representative snapshots of experimental plants grown under each condition (i.e., bare soil
252 or soil containing SHS and N or R irrigation) at different weeks during their growth, from
253 transplanting (week 0) to fruit ripening (week 12), are shown in **Fig. 1**. Overall, the plants
254 grown in soils containing SHS were larger and appeared healthier than those grown in bare
255 soil.

256



257

258

259 **Fig. 1. Effects of SHS mulching on plants.** (A) Schematics of the evaporation and transpiration pathways
260 and their impacts on plants. (B) Representative photographs of tomato plants grown in controlled growth
261 chambers mulched with 1-cm-thick superhydrophobic sand (SHS) and bare soil plants under normal (N) and
262 reduced (R) irrigation. Tomato plants were transplanted and grown for 98 days until the final fruit and
263 biomass harvest. Each treatment combination involved four ($n = 4$) replicate plants.

264

265 **Soil water content**

266 The daily SWC corresponding to each irrigation level is detailed in **Fig. 2A**. For both N
267 and R irrigation strategies (i.e., 100% and 50% field capacity, respectively), the SWC was
268 slightly increased during days 40–98 to compensate for the increase in plant size and water
269 demand. The slight fluctuations in SWC below the initial field capacities observed at five
270 occasions (i.e., between 30–50 days) represent five days when pot weights were recorded
271 without adding water until the following day due to unavoidable circumstances. However,
272 the plants were not severely stressed on these occasions, as the available SWC was enough
273 to meet the ET demand of the plants until the next day when water was added to the pots.
274 When averaged across both mulched and bare soils, the mean SWC was 0.62 ± 0.01 and
275 0.33 ± 0.01 kg/pot under N and R irrigation, respectively. This result also demonstrates
276 that plant growth could be maintained under 50% of normal irrigation (i.e., the R irrigation
277 strategy).

278

279 **Evapotranspiration**

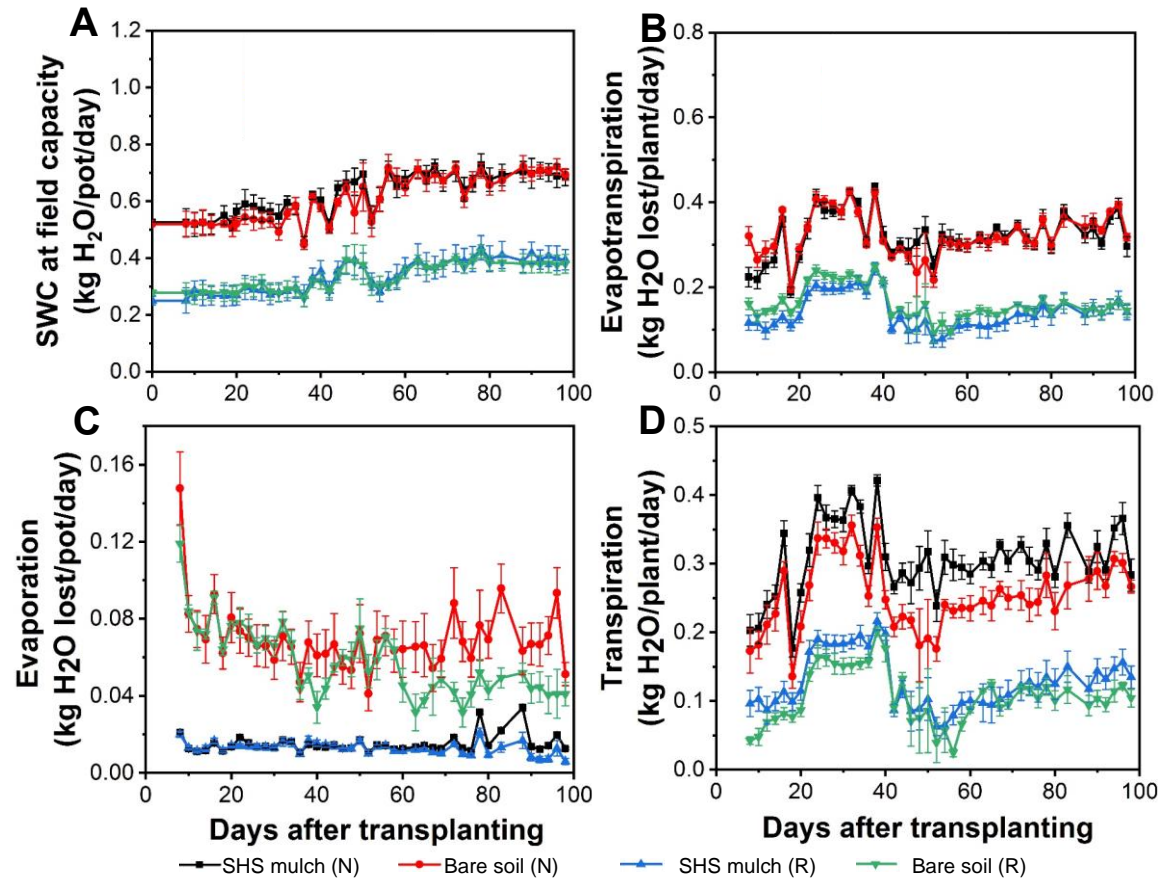
280 The daily ET significantly differed between the N and R irrigation scenarios and accounted
281 for approximately 50% of the daily SWC on average, as shown in **Fig. 2B**. Under R
282 irrigation, ET differed between mulched and bare soils, notably during days 10–35 and 45–
283 75; no significant differences were found under N irrigation.

284

285 **Soil evaporation and transpiration**

286 Throughout the growth period, the daily evaporation was significantly lower from soil
287 containing SHS than bare soil under N and R irrigation, as shown in **Fig. 2C**. Consequently,
288 plants in mulched soil had higher daily transpiration than in bare soil under both irrigation
289 scenarios, as shown in **Fig. 2D**.

290



291

292

293 **Fig. 2.** (A) Soil water content (SWC) in pots maintained at field capacity for each treatment and irrigation
294 regime, (B) daily changes in ET, (C) evaporation from the soil, and (D) transpiration between SHS-mulched
295 and bare soils under N and R irrigation strategies throughout plant growth. Each data point is a mean of four
296 ($n = 4$) replicates; error bars are standard errors (\pm SE) of the means.

297

298 Cumulative evapotranspiration

299 There was no significant difference in the cumulative ET loss between mulched and bare
300 soils under N irrigation; both had a mean cumulative ET of 13.9 ± 0.7 kg/plant over the 98
301 days of observation, as shown in **Fig. 3A**. Under R irrigation, the cumulative ET was 13%
302 higher in SHS-mulched soil in than bare soils (with mean values of 6.97 ± 0.57 and $6.06 \pm$
303 0.78 kg/plant, respectively).

304

305

306 **Cumulative soil evaporation and transpiration**

307 In SHS-mulched soils, that total cumulative evaporation (and transpiration) accounted for
308 5% (95%) and 9% (91%) of the total ET under **N** and **R** irrigation, respectively; in bare
309 soils, evaporation (and transpiration) accounted for 21.5% (78.5%) and 31.5% (65.5%) of
310 the total ET under **N** and **R** irrigation, respectively (**Figs. 3B and 3C**). Overall, SHS
311 mulching suppressed evaporation and enhanced transpiration by the same magnitude of
312 78% and 17%, respectively under both **N** and **R** irrigation relative to the bare soils.

313

314 **Plant height**

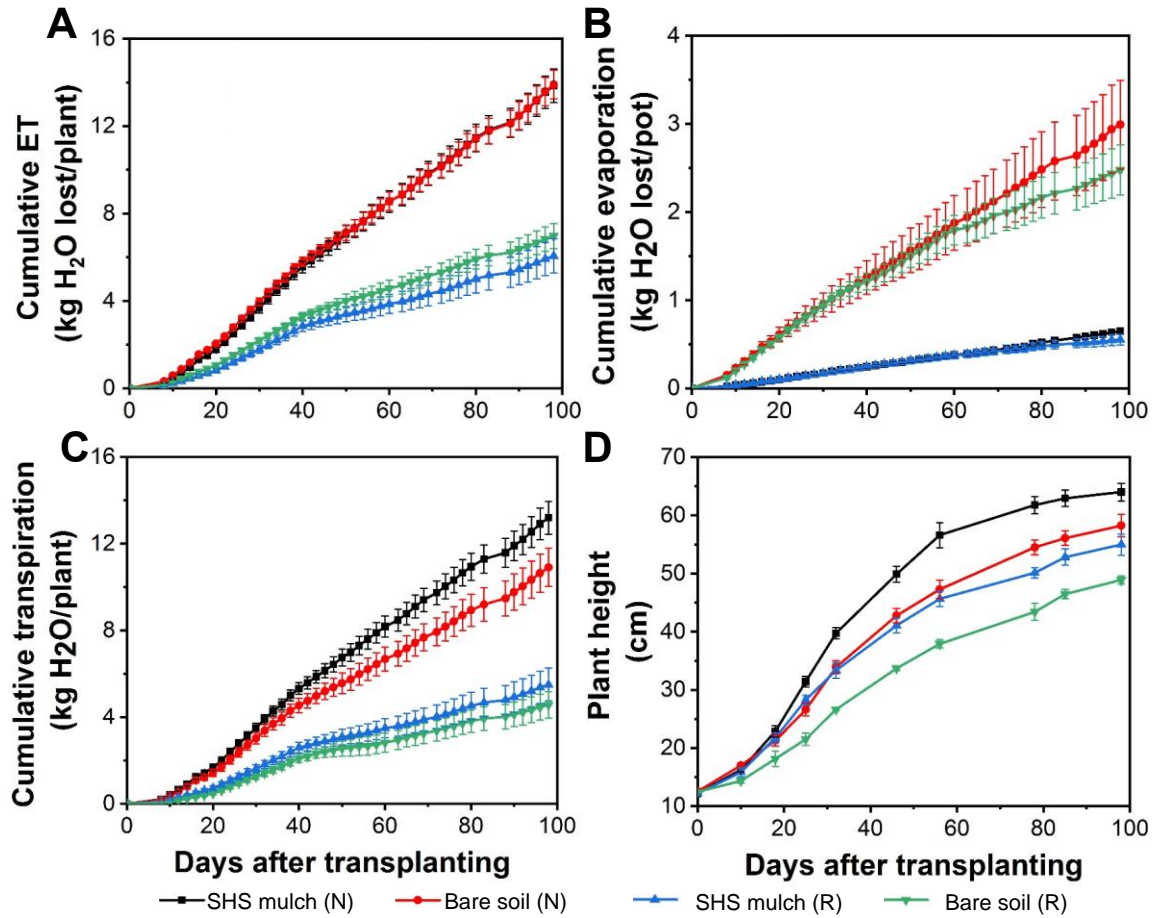
315 Throughout the growth period, the maximum plant height in SHS-mulched soil
316 significantly exceeded that in bare soil by 9% ($p = 0.005$) and 11% ($p < 0.05$) under **N** and
317 **R** irrigation, respectively (**Fig. 3D**). In particular, a more rapid increase in the plant height
318 was observed during 10–45 days after transplanting; after 70 days, plant height started to
319 level off.

320

321 **Dependence of evapotranspiration on plant growth stage**

322 The daily mean ET, evaporation, and transpiration during days 0–30, 31–60, and 61–98
323 are compiled in **Table 1**. Overall, the mean ET, evaporation, and transpiration increased
324 with time; significant differences were present in the mean ET and transpiration between
325 each timeframe ($p < 0.05$). However, no significant difference was found in the mean
326 evaporation from bare soil with time under **N** irrigation ($p > 0.05$), as evidenced by the
327 three-way ANOVA results. Significant two-way interactions were present between
328 mulching and irrigation regimes on transpiration ($p < 0.05$) and between irrigation regime
329 and growth stage on ET, evaporation, and transpiration ($p < 0.05$).

330



331

332

333 **Fig. 3.** (A) Cumulative evapotranspiration (ET), (B) evaporation, (C) transpiration, and (D) plant height
334 during plant growth under contrasting soil mulching treatments (i.e., SHS-mulched vs. bare soil) and
335 irrigation regimes (i.e., N and R). Each data point is a mean of four ($n = 4$) replicates; error bars represent
336 the \pm SE of the mean.

337 **Table 1. Effects of soil mulching (M), irrigation regimes (W), growth stage (S)– shown by the three different time periods, and**
 338 **their interactions on mean evapotranspiration (ET), evaporation, and transpiration.**

Treatments	ET (kg/plant)			Evaporation (kg/pot)			Transpiration (kg/plant)		
	0–30 days	31–60 days	61–98days	0–30 days	31–60 days	61–98 days	0–30 days	31–60 days	61–98 days
SHS mulch (N)	3.66±0.14	4.89±0.21	5.29±0.22	0.19±0.02	0.21±0.01	0.27±0.01	3.48±0.13	4.68±0.20	5.02±0.22
Bare soil (N)	3.93±0.14	4.59±0.16	5.34±0.15	0.93±0.11	0.93±0.16	1.12±0.18	3.00±0.21	3.66±0.20	4.23±0.30
SHS mulch (R)	1.79±0.14	2.07±0.24	2.20±0.33	0.18±0.40	0.20±0.02	0.18±0.02	1.61±0.13	1.86±0.23	2.02±0.31
Bare soil (R)	2.24±0.15	2.39±0.17	2.40±0.11	0.92±0.02	0.86±0.08	0.68±0.01	1.32±0.15	1.57±0.22	1.74±0.07
3-way ANOVA	ET			Evaporation			Transpiration		
M (df =1)	ns			266 (0.0*)			25.0 (0.0*)		
W (df =1)	702 (0.0*)			5.63 (0.02)			500 (0.0*)		
S (df = 2)	33.0 (0.0*)			ns			26.0 (0.0*)		
M×W (df = 1)	ns			ns			5.28 (0.03)		
M×S (df = 2)	ns			ns			ns		
W×S (df = 2)	15.0 (0.0*)			3.49 (0.04)			7.74 (0.0*)		
M×W×S (df = 2)	ns			ns			ns		

339 Data presented in the upper rows indicate mean values (\pm SE) of total ET, evaporation, and transpiration at three time periods (0–30, 31–60, and 61–98 days after
 340 transplanting) for each treatment combination: SHS-mulched mulch under N irrigation (SHS mulch-N); bare soil under N irrigation (Bare soil-N); SHS-mulched
 341 mulch under R irrigation (SHS mulch-R); and bare soil under R irrigation (Bare soil-R). The lower rows indicate three-way ANOVA statistics for the effects of
 342 M, W, S, and their interactions on total ET, evaporation, and transpiration. Significant results are represented by *F* values followed by their respective *p* (*p*
 343 < 0.05) in parentheses; ns = non-significant results; 0.0* represents *p*-values < 0.001; df = degrees of freedom.

344 **Effects of SHS mulching on physiological traits, biomass yield, and fruit yield**

345 The performed two-way ANOVA demonstrated the significant effect of SHS mulching on
 346 total evaporation and transpiration ($p < 0.05$) but no significant effect of SHS mulching on
 347 the total ET, as presented in **Table 2**. In addition to the changes in ET, evaporation,
 348 transpiration, and plant height demonstrated in **Figs. 2–3**, SHS mulching impacted other
 349 plant responses such as stomatal conductance and aperture, CCI, TE, fruit yields, HI, fresh
 350 mass, dry mass, root xylem vessel, and total root diameter, as presented in **Figs. 4–7**.

351

352 **Table 2. Results of two-way ANOVA test showing effects of soil mulching (M),**
 353 **irrigation regime (W), and their interactions (M × W) on plant traits measured.**

Source of variation	df	Total ET	Total evaporation	Total transpiration	Stomatal conductance	Chlorophyll content index, CCI
M	1	ns	92.7 (0.0*)	12.3 (0.0*)	17.5 (0.0*)	62.6 (0.0*)
W	1	197 (0.0*)	ns	232 (0.0*)	23.5 (0.0*)	17.4 (0.0*)
M×W	1	ns	ns	ns	ns	ns
Error		0.002	0.196	0.849	15810	26.584

Source of variation	df	Plant height	Shoot fresh mass	Root fresh mass	Total fresh mass	Shoot dry mass
M	1	11.7 (0.01)	51.4 (0.0*)	15.8 (0.0*)	53.2 (0.0*)	21.5 (0.0*)
W	1	43.4 (0.0*)	89.9 (0.0*)	29.9 (0.0*)	111 (0.0*)	80.3 (0.0*)
M×W	1	ns	12.5 (0.0*)	ns	11.2 (0.0*)	7.65 (0.02)
Error		6.468	103.310	5.971	114.8	5.619

Source of variation	df	Root dry mass	Total dry mass	Transpiration efficiency, TE	Fruit yields per plant	Harvest Index, HI
M	1	20.6 (0.0*)	14.1 (0.0*)	ns	6.19 (0.03)	15.6 (0.0*)
W	1	ns	36.2 (0.0*)	45.1 (0.0*)	23.3 (0.0*)	12.2 (0.0*)
M×W	1	ns	ns	ns	ns	ns
Error		0.792	10.518	0.1326	1541	0.004

354 Data presented are *F* values from two-way ANOVA and p-values for significant results ($p < 0.05$) in
 355 parentheses; ns = non-significant results. 0.0* represents p -values < 0.001 ; df = degrees of freedom.

356

357

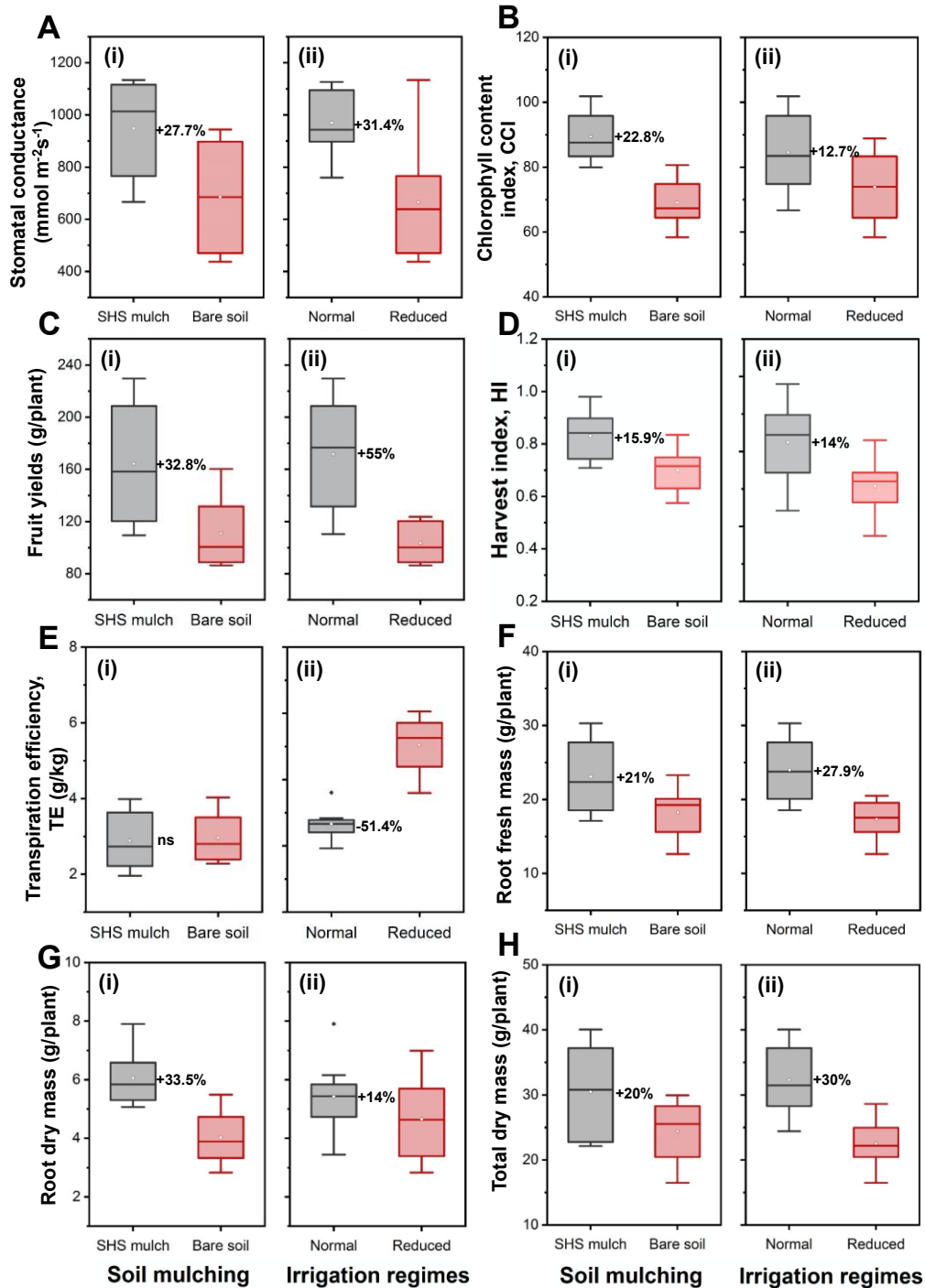
358 **Main effects of soil mulching and irrigation regimes**

359 A 27.7% higher stomatal conductance was present in SHS-mulched plants than those in
360 bare soil across irrigation regimes ($p = 0.001$; **Fig. 4A-i**). Stomatal conductance was also
361 31.4% higher under **N** irrigation than under **R** irrigation ($p < 0.05$; **Fig. 4A-ii**). Similarly,
362 leaf CCI was 22.8% higher in SHS-mulched plants than those in bare soil ($p < 0.05$; **Fig.**
363 **4B-i**). Irrespective of mulch treatment, leaf CCI was 12.7% higher under **N** irrigation than
364 **R** irrigation ($p = 0.001$; **Fig. 4B-ii**). Fruit yield, defined by the total fruit mass per plant,
365 was 32.8% higher in mulched soil than bare soil ($p = 0.03$; **Fig. 4C-i**) and 55% higher
366 under **N** irrigation than **R** irrigation ($p < 0.05$; **Fig. 4C-ii**).

367 The relationship between fruit yield and total fresh mass (i.e., the HI) was
368 characterized by 16% higher HI in mulched soil than bare soil ($p = 0.002$; **Fig. 4D-i**).
369 Regardless of the mulch treatment, using **R** irrigation decreased the HI by 14% ($p = 0.004$;
370 **Fig. 4D-ii**). No significant difference in the ratio of shoot dry mass to total water transpired
371 per plant (i.e., the TE) was present between plants grown in mulched and bare soils ($p =$
372 0.697 ; **Fig. 4E-i**). However, the TE was 51.4% lower under **N** irrigation than **R** irrigation
373 ($p < 0.05$; **Fig. 4E-ii**).

374 In terms of biomass allocation, plants in mulched soil had 21% more root fresh
375 mass than those in bare soil ($p = 0.002$; **Fig. 4F-i**); plants under **N** irrigation had 28% more
376 root fresh mass than their **R** irrigation counterparts ($p < 0.05$; **Fig. 4F-ii**). The root dry mass
377 was 33.5% higher in mulched soil than bare soil ($p < 0.05$; **Fig. 4G-i**), while it was 14%
378 higher under **N** irrigation than **R** irrigation ($p = 0.046$; **Fig. 4G-ii**). The total dry mass in
379 mulched soil increased by 20% over that in bare soil ($p = 0.003$; **Fig. 4H-i**), while **N**
380 irrigation resulted in 30% higher total dry mass than under **R** irrigation ($p < 0.05$; **Fig. 4H-**
381 **ii**).

382



383

384

385 **Fig. 4.** ANOVA results showing effects of (i) soil mulching (SHS mulch vs. bare soil) and (ii) irrigation

386 regimes (N vs. R) on (A) leaf stomatal conductance; (B) CCI; (C) fruit yields per plant; (D) HI; (E) TE; (F)

387 root fresh mass; (G) root dry mass; and (H) total dry mass per plant. Each box represents the data distribution

388 from eight plants ($n = 8$) with the median along the mid-line. The white dot inside the box represents the

389 mean value, the upper and lower sections of the box represent the 25% and 75% confidence intervals,

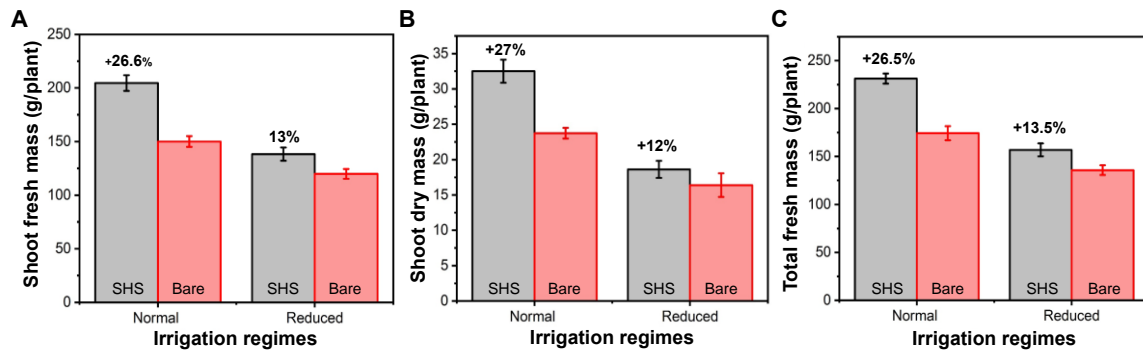
390 respectively, the whiskers on the box represent the 1.5 interquartile range, and dots outside the box indicate
391 outliers. ns: no significant difference between treatments.

392

393 **Interaction effects of mulching and irrigation regimes**

394 There was also a significant interaction effect of mulching and irrigation on shoot fresh
395 mass, shoot dry mass, and total fresh mass per plant. The shoot fresh mass was 26.6%
396 higher in mulched soil than bare soil under **N** irrigation, whereas under **R** irrigation, the
397 shoot fresh mass was 13% higher in mulched soil than bare soil (**Fig. 5A**; $p < 0.05$, $p =$
398 0.019 , respectively). The shoot dry mass increased in mulched soil by 27% compared with
399 that in bare soil under **N** irrigation; under **R** irrigation, shoot dry mass was 12% higher in
400 mulched soil than bare soil (**Fig. 5B**; $p = 0.001$, $p = 0.046$, respectively). Under **N**
401 irrigation, SHS mulching increased the total fresh mass per plant by 26.5%; under **R**
402 irrigation, the total fresh mass per plant was 13.5% higher in mulched soil than bare soil
403 (**Fig. 5C**; $p < 0.05$, $p = 0.037$, respectively).

404



405

406

407 **Fig. 5.** Two-way interaction effects of soil mulching (SHS vs. bare soil) and irrigation regimes (**N** vs. **R**) on
408 (A) shoot fresh mass, (B) shoot dry mass, and (C) total fresh mass per plant. Each bar represents the mean
409 value of four replicates ($n = 4$), and each error bar represents the SE of the means of four samples.

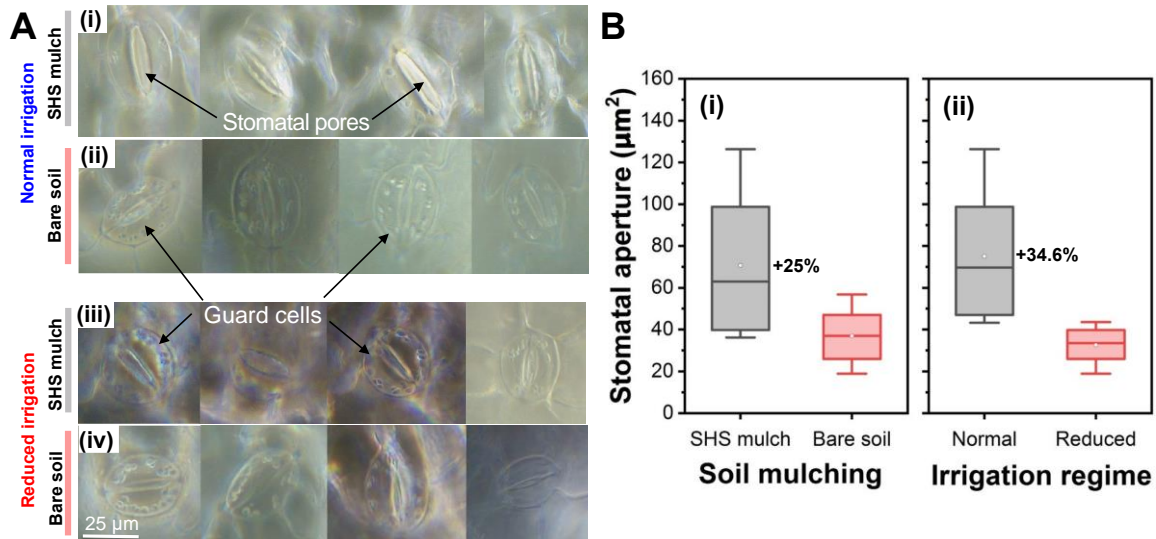
410

411 **Leaf stomatal status**

412 Microscopy revealed that the stomatal pores of plants grown in mulched soils (**Fig. 6A-i**
413 **& iii**) were larger than those in bare soils (**Fig. 6A-ii & iv**); the mean stomatal aperture
414 (area) was 25% larger regardless of the irrigation regime ($p = 0.0019$; **Fig. 6B-i**).

415 Furthermore, the mean stomatal aperture was 34.6% higher under **N** irrigation than **R**
416 irrigation ($p < 0.05$; **Fig. 6B-ii**).

417



418

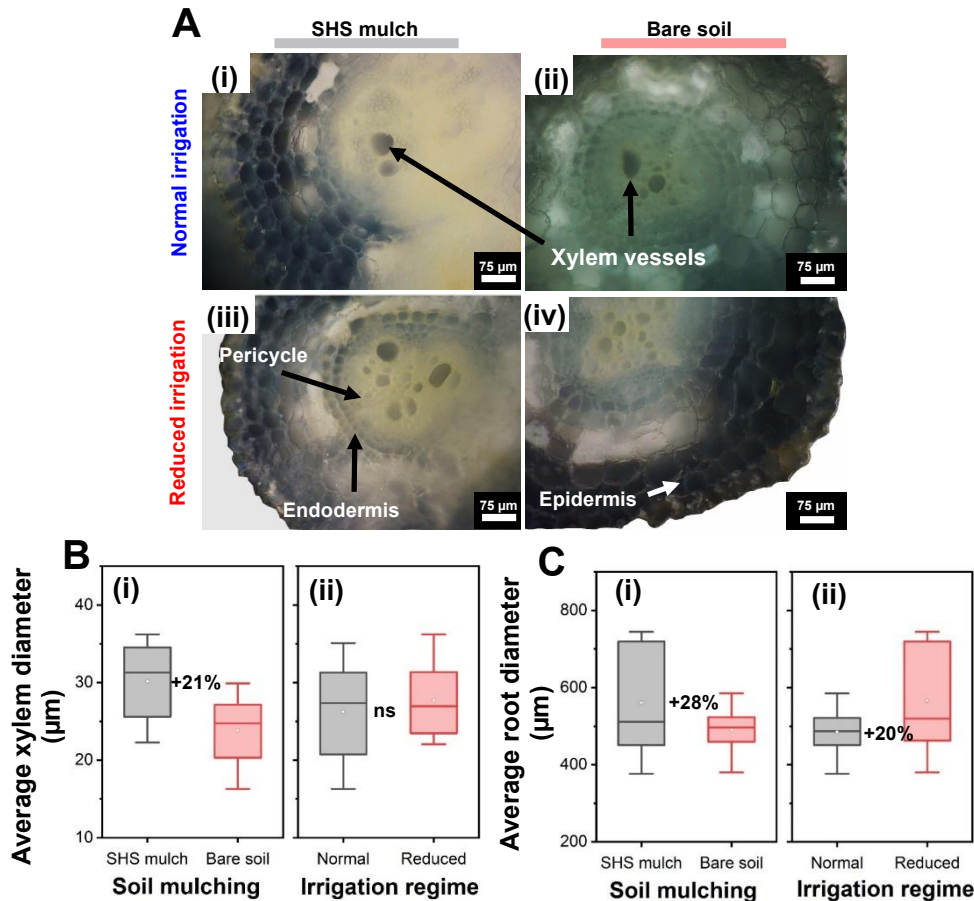
419

420 **Fig. 6.** Microscopic analysis of leaf stomatal pores in response to soil mulching and irrigation regimes as on
421 the final date of plant harvest. **(A)** Leica DVM6 microscopic image of leaf stomata for plants grown under **N**
422 and **R** irrigation in **(i, iii)** SHS-mulched and **(ii, iv)** bare soils. **(B)** Boxplots showing the mean stomatal
423 aperture (area) for plants grown in **(i)** SHS and bare soil, and **(ii)** under **N** and **R** irrigation. Each box
424 represents the data distribution from 16 leaf samples ($n = 16$) with the median along the mid-line. The white
425 dot inside the box represents the mean value, the upper and lower sections of the box represent the 25% and
426 75% confidence intervals, respectively, and the whiskers on the box represent the 1.5 interquartile range.

427

428 **Root anatomical structure**

429 Finally, root anatomical features were measured to determine the shoot–root feedback
430 linkages with the observed trends in ET as a function of mulching and irrigation scenarios;
431 these results are summarized in **Figs. 7A-i–7A-iv**. The xylem vessel diameter was 21%
432 larger for roots grown in mulched soil than bare soil ($p = 0.001$; **Fig. 7B-i**). However, the
433 difference in diameter under differing irrigation scenarios was not statistically significant
434 ($p = 0.343$; **Fig. 7B-ii**). Additionally, the root diameter increased in mulched soil by 28%
435 relative to bare soil ($p = 0.0177$; **Fig. 7C-i**), whereas it decreased from **R** irrigation to **N**
436 irrigation by 20% ($p = 0.0067$; **Fig. 7C-ii**).



437

438

439 **Fig. 7.** (A, i-iv) Cross-sections of tomato roots from contrasting soil mulching and irrigation scenarios. (B-

440 C) Box plots showing the effects of (i) SHS mulching and (ii) irrigation regimes on (B) average xylem vessel

441 diameter and (C) average root diameter. Each box represents the data distribution from 16 root samples ($n =$

442 16) with the median along the mid-line. The white dot inside the box represents the mean value, the upper

443 and lower sections of the box represent the 25% and 75% confidence intervals, respectively, the whiskers on

444 the box represent the 1.5 interquartile range, and dots outside the box indicate outliers. ns: no significant

445 difference between treatments.

446

447 DISCUSSION

448 To clarify the effects of SHS mulching presented on ET fluxes and phenotypic traits of

449 tomato plants grown under N and R irrigation, the mechanistic insights behind SHS

450 mulching and their various impacts on plants and interrelationships are analyzed in this

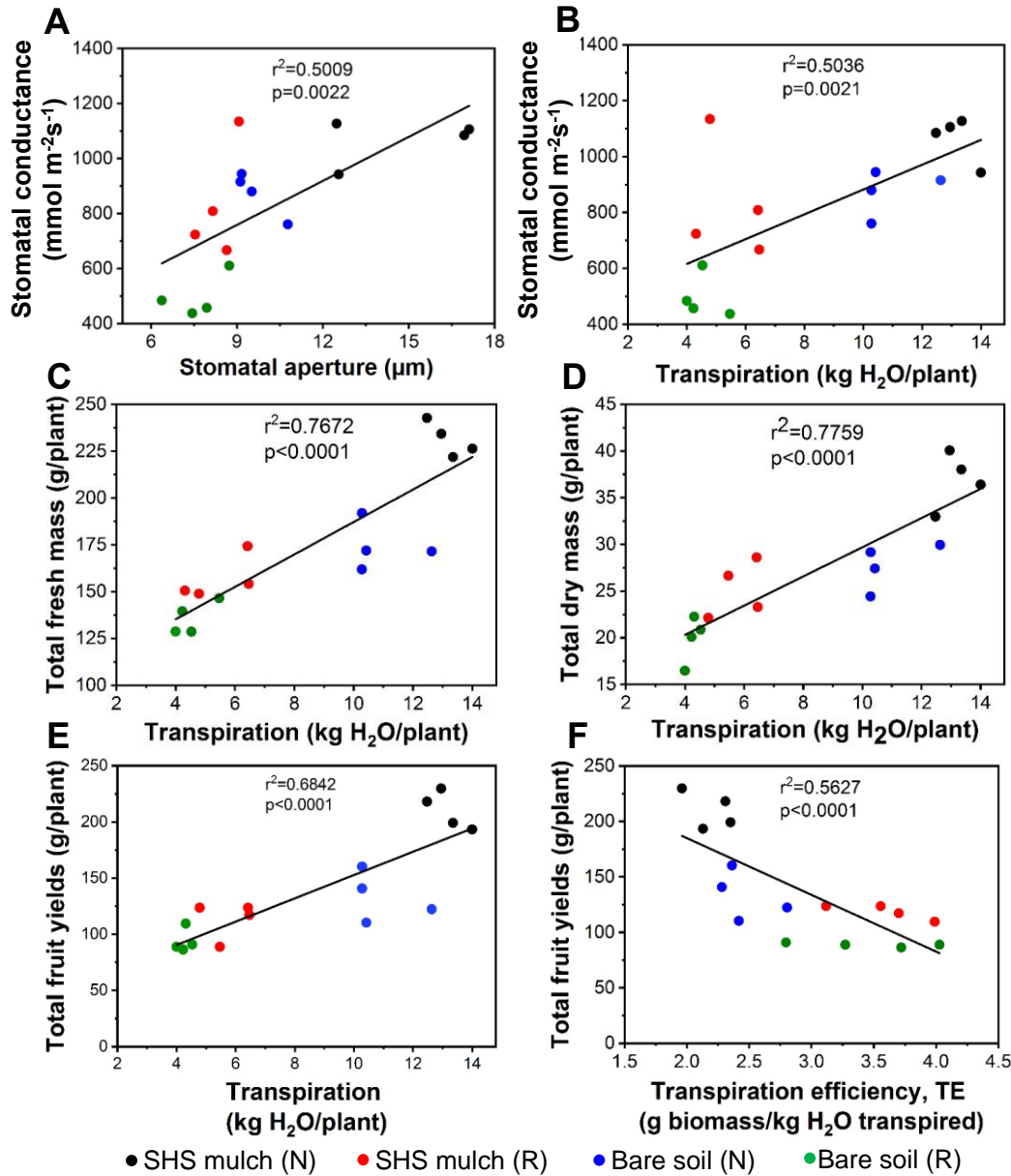
451 section. The physical mechanism behind the ability of SHS to reduce evaporation from soil

452 is based on the Laplace pressure (i.e., the pressure difference across a curved liquid

453 surface), which depends on the surface tension of water and the curvature of the liquid–air
454 interface (Domingues et al., 2018, Das et al., 2019, Gallo Jr et al., 2021a). When water is
455 supplied through sub-surface irrigation, it rises upward through the soil by capillarity and
456 contacts the SHS layer on the topsoil. The curvature of the air–water interface touching the
457 SHS grains forms a convex shape; the capillary action that drives the rise of water in soils
458 prevents it from wetting the SHS layer, creating a diffusion barrier (Gallo Jr et al., 2021a,
459 Gallo Jr et al., 2021b). By physically covering the topsoil, the mulch also insulates the soil
460 from solar radiation and wind, thereby further reducing the capacity for evaporation losses
461 (Kader et al., 2017, Gallo Jr et al., 2021a). With less evaporation, the soil moisture content
462 rises, allowing for higher plant transpiration.

463 A significant causal relationship was present between the stomatal conductance and
464 aperture (**Fig. 8A**) and between transpiration and stomatal conductance (**Fig. 8B**), total
465 fresh mass (**Fig. 8C**), total dry mass (**Fig. 8D**), and fruit yield (**Fig. 8E**). These relationships
466 suggest that the soil moisture-retaining function of SHS augments a chain of plant-related
467 responses that affect stomatal opening and closure, which directly influence transpiration
468 rates. Plants can alter their stomatal pore aperture by actively adjusting the turgor pressure
469 within their guard cells (Blatt, 2000), which moderates the exchange of gases such as water
470 vapor, carbon dioxide (CO₂), and oxygen between the leaf interior and the atmosphere
471 (Kollist et al., 2014, Lawson and Vialet-Chabrand, 2019, Franks and Farquhar, 2007).
472 Enhanced water availability in the soil increases the guard cell turgor pressure, leading to
473 a larger stomatal aperture and increased stomatal conductance of water vapor, which
474 enhances the transpiration and CO₂ uptake rates for photosynthesis (Bertolino et al., 2019,
475 Condon et al., 2002, Putra et al., 2012). This rationale underpins the observed increase in
476 transpiration (**Figs. 2–3; Table 1**), stomatal conductance (**Fig. 4A**), and stomatal aperture
477 (**Fig. 6**) when using SHS mulch.

478



479

480 **Fig. 8.** Regression analyses showing the relationship between (A) leaf stomatal conductance and stomatal
 481 aperture, (B) stomatal conductance and total transpiration, (C) total fresh mass and total transpiration, (D)
 482 total dry mass and total transpiration, (E) total fruit yield per plant and total transpiration, and (F) total fruit
 483 yield per plant and TE. Each dot on the graph indicates an individual plant grown in SHS mulch or bare soil
 484 under N or R irrigation.

485

486 Aboveground physiological processes, including stomatal responses, are governed
 487 by a feedback mechanism linked closely with the hydraulic conductivity of the soil water

488 in the root zone (Putra et al., 2012, Tardieu and Parent, 2017, Zarebanadkouki et al., 2016,
489 Bertolino et al., 2019, Adachi et al., 2010, Chaves et al., 2002). This hydraulic conductivity
490 is dependent on the diameter of the roots and the xylem vessels (Niklas, 1985, Comas et
491 al., 2013). Here, larger xylem vessel and root diameters were present in mulched soil than
492 in bare soil (**Fig. 7**). Consistent with these findings, larger xylem vessel and root diameters
493 have been shown to develop in wheat plants in response to increased SWC caused by
494 mulching (Zhan et al., 2019), increasing WUE and shoot biomass (Kadam et al., 2015).
495 Although small-diameter roots maintain plant growth during a drought, especially at soil
496 depths with available water (Comas et al., 2013), several researchers have suggested that
497 large-diameter roots are more instrumental in resource uptake and plant growth under water
498 scarcity (Larson and Funk, 2016, Kadam et al., 2015). The latter agrees with the observed
499 results; the roots had a larger average diameter under **R** irrigation than **N** irrigation. A larger
500 diameter may promote root penetrative ability or soil exploration in dry soils (Clark et al.,
501 2003) and enhance root longevity (McCormack et al., 2012). These results indicate the
502 important role that root architecture plays in resource acquisition and conservation to
503 facilitate aboveground processes that determine productivity in plants.

504 The root–shoot hydraulic system conducts nutrients and water via the transpiration
505 stream to the leaves; these nutrients (e.g., nitrogen) support plant growth and the
506 photosynthetic process. Nitrogen availability is related to the total chlorophyll content in
507 plant leaves (Bassi et al., 2018). The higher the chlorophyll content, the more efficient its
508 photosynthetic capacity is, causing increased growth, yields, and biomass (Condon et al.,
509 2004, Haeferle et al., 2009). Here, a higher CCI and higher transpiration due to SHS
510 mulching contributed to higher fruit yield and biomass production, leading to a higher HI.
511 The fruit mass per plant was negatively correlated with TE (**Fig. 8F**), indicating that fruit
512 yields increased under low TE. This finding contradicts the proposal that higher TE is
513 critical in increasing crop yields in water-limited environments (Christy et al., 2018,
514 Sinclair, 2012, Rebetzke et al., 2002, Coupel-Ledru et al., 2016, Condon et al., 2004, Vadez
515 and Ratnakumar, 2016). However, TE is a physiologically complex trait that results from
516 a combination of multiple physiological parameters, such as photosynthesis, stomatal
517 conductance, mesophyll conductance (i.e., the diffusion of CO₂ from sub-stomatal cavities
518 to the sites of carboxylation in the chloroplasts) (Flexas and Medrano, 2002, Flexas et al.,

519 2008), and other conditions that determine carbon balance and growth in plants (Natarajan
520 et al., 2021). Consequently, the conditions responsible for the expression of high TE in one
521 situation may be associated with a low TE in another environment (Sinclair, 2012).

522 A prolonged stomatal opening and higher stomatal conductance account for lower
523 TE (Flexas et al., 2008); mulched soils retain more SWC and maintain stomatal openings
524 longer than bare soils, leading to a low TE (Balwinder-Singh et al., 2011). However, no
525 significant difference in TE was present between SHS-mulched and bare soils, despite their
526 difference in stomatal conductance. However, lower stomatal conductance was associated
527 with higher TE under **R** irrigation than under **N** irrigation. These findings indicate the trade-
528 off required between stomatal conductance and carboxylation efficiency to improve TE
529 and yields. Unlike increasing the stomatal conductance, increasing the mesophyll
530 conductance increases photosynthesis and TE without increasing transpiration because the
531 CO₂ diffusion pathway involving mesophyll conductance is not shared with that of
532 transpired water (Flexas et al., 2008, Ouyang et al., 2017, Flexas et al., 2013, Barbour et
533 al., 2010).

534 Future investigations are needed to examine physiological traits associated with
535 enhanced TE under different environmental conditions. For example, experiments
536 resolving instantaneous leaf gas exchange could help determine photosynthesis and
537 conductance (stomatal vs. mesophyll conductance), whereas measurements of carbon
538 isotope composition ($\delta^{13}\text{C}$; a surrogate trait that integrates TE and variation in root traits
539 and plasticity) could enhance understanding of how aboveground and belowground
540 processes shape plant responses and yield performance under different water regimes.
541 Determining the genetic factors that can maximize transpiration and carbon assimilation to
542 enhance photosynthetic output would complement the water-saving benefit of SHS
543 mulching to improve yields in arid land agriculture.

544

545 **CONCLUSIONS**

546 In conclusion, by combining a broad set of techniques to investigate the effects of SHS
547 mulching on ET fluxes and plant responses, this study revealed that adding a thin (1.0-cm
548 thick) layer of SHS dramatically decreased water evaporation from the topsoil, thereby
549 boosting transpiration under both irrigation scenarios studied. The enhanced transpiration,

550 in turn, increased plant biomass and fruit yield. Furthermore, this study facilitated
551 understanding the mechanistic insights behind increased transpiration, biomass, and fruit
552 yield owing to the root–shoot hydraulic processes accounting for enhanced water uptake
553 and stomatal regulation. The quantitative assessment of ET, evaporation, and transpiration
554 presented here will aid crop WUE assessments and field water optimization and
555 management using SHS mulching, irrigation systems design, and agricultural regimes in
556 arid and semi-arid regions.

557

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561

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565

566 **AUTHOR CONTRIBUTIONS**

567 **KO:** Designed and performed the experiments, including soil preparation, planting,
568 harvests, data collection, and analysis; manuscript writing and revision. **AGJ:** Assisted
569 with the experimental design, data analysis, and manuscript revision. **VD:** Collected the
570 evapotranspiration data and assisted in plant harvest and biomass determination. **HM:**
571 Supervised this research and revised the manuscript.

572

573 **DATA AVAILABILITY STATEMENT**

574 All data supporting the findings reported in this study are available in the paper. **HM** and
575 **AGJ** have filed a US patent (US20200253138A1) for the SHS used in this work, while
576 **KO** and **VD** have no competing financial interest in this work.

577

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