### Quantifying river water contributions to riparian trees along a losing river: Lessons from stable isotopes and iteration method

Yue Li<sup>1, 2</sup>, Ying Ma<sup>1, 2</sup>, Xianfang Song<sup>1, 2</sup>, Qian Zhang<sup>3</sup>, Lixin Wang<sup>4</sup>

<sup>1</sup>Key Laboratory of Water Cycle and Related Land Surface Processes, Institute of Geographic Sciences and Natural Resources Research, Chinese Academy of Sciences, Beijing 100101, China

<sup>2</sup>University of Chinese Academy of Sciences, Beijing 100049, China

<sup>3</sup>Institute of Geographic Sciences and Natural Resources Research, Chinese Academy of Sciences, Beijing 100101, China

<sup>4</sup>Department of Earth Sciences, Indiana University-Purdue University Indianapolis (IUPUI), Indianapolis, IN 46202, United States

Correspondence to: Ying Ma (maying@igsnrr.ac.cn)

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Abstract. River water plays a critical role in riparian plant water use and riparian ecosystem restoration along losing rivers (rivers losing flow into underlying groundwater) under warming climates warming. How to quantify 15 the contributions of river water to riparian plants under different water tablegroundwater levelss and the related responses of plant water use efficiency is a great challenge. In this study, experiments observations of stable water isotopes ( $\delta^2 H_{-}$  and  $\delta^{18}O_{-}$  and  $\delta^{18}O_{-}$  and  $\delta^{222}Rn$ ), 222Rn, in different waters and leaf  $\delta^{13}C$  were conducted for riparian-deeprooted riparian weeping willow (Salix babylonica (L.) during 2019 dry year (-dry year-2019) and 2021 wet year (wet year2021) along the Chaobai River in Beijing, China. We The MixSIAR model in combination with 20 anproposed an -iteration method in combination with the MixSIAR model were proposed to quantify the proportional river water contribution (RWC) to transpiration flux-of riparian S. babylonica and its correlations with the depth of to the water table water table depth (WTD) and leaf  $\delta^{13}$ C. Results showed that riparian S. *babylonica* took up deep water (in the 80-170 cm soil layer and groundwater) by  $56.5 \pm 10.8\%$ . River water that recharged riparian deep water was an indirect water source and contributed by by 20.3% of water to riparian trees 25 nearby the losing river. Significantly increasing river water acquisitions uptake (by 7.0%) but and decreasing leaf  $\delta^{13}$ C (by -2.0%) of riparian trees were observed as the WTD-water table depth changed from 2.7 m in dry 2019 to 1.7 m in wet 2021 (p < 0.05).- The higher water availability probably promoted stomatal opening and thus increasing transpiration water loss, which led to the decreasing leaf  $\delta^{13}$ C in wet year compared to dry year. A short residence time (no more than 0.28 days) of groundwater indicated that there was rapid and frequent river recharge

- 30 to riparian groundwater in 2021. It was found that the RWC-river water contribution to riparian *S. babylonica* was negatively correlated with the WTD-water table depth but positively related to theand leaf  $\delta^{13}$ C in linear functions (p = < 0.000001). The rising-water table groundwater level would stimulate trigger riparian trees to increase the water extraction from maximize groundwater/river transpiration water consumptions and show a- consumptive river-water-use pattern, which could not be recommended in order to protect both rivers and riparian
- 35 vegetation.profligate water use strategy with increasing water extraction from the losing river. This study provides critical insights into understanding the mechanisms of water cycle in Groundwatergroundwater-Soilsoil-Plantplant-Atmosphere\_atmosphere\_Continuum, managing water resources and riparian afforestation along losing rivers.

#### **1** Introduction

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- Ongoing climate warming as well asand groundwater overexploitation hashave altered precipitation regimes, river flow-runoff and bank storage globally, further leading to widespread riskse of rivers losing flow into underlying groundwater ("losing" river) and even running dry (Winter et al., 1998; Schindler and Donahue, 2006; Allen et al., 2015; Jasechko et al., 2021). Ecological wWater replenishment of to losing rivers and riparian revegetation have been pushed forward worldwide to restore the river ecosystem (Smith et al., 2018; Long et al., 2020). The Wwater replenishing-replenishment to losing rivers contributed byby \_40% to bank storage and groundwater storage recovery (Long et al., 2020). However, large-scale riparian revegetation increased the\_plant transpiration substantially, which in turn led to great loss of riparian bank storage and even river flow-runoff (Moore and Owens, 2012; Dzikiti et al., 2013; Missik et al., 2019; Mkunyana et al., 2019). Therefore, deeply understanddetermining what water sourceshere and how much river water is taken up by riparian trees-took up and their responses of tree
- 50 <u>water use characteristics</u> to the variations in the groundwater levelwater table variations could can help to implement management strategies for maintaining balance the river flow runoff and tree water requirement need of revegetated riparian species zones.

The potential water sources of riparian trees along a losing river are generally considered as a mix of soil water at different depths, groundwater, and river water (Alstad et al., 1999; White and Smith, 2020). However, there wais a drastic debate on whether the river water is a potential water source for of riparian trees or not and how it becaomes available to plants. The river water contribution (RWC) to riparian trees has been widely estimated using the data comparison, graphical inference, two-or multi-source linear mixing models and Bayesian mixing models (MixSIR, SIAR, SISUS, MixSIAR) accompanied with stable water isotopes (8<sup>2</sup>H and 8<sup>44</sup>O) (Dawson and Ehleringer, 1991; Ehleringer and Dawson, 1992; White and Smith, 2020).-Most of previous studies considered
the river water as a separate-direct water source to quantify-evaluate the river water contribution (RWC) to transpiration fluxuptake of riparian trees (Alstad et al., 1999; Zhou et al., 2017; White and Smith, 2020). A number of previous These studies showed that the separate the river water directly source contributed up to 80% to riparian plant transpiration directly-based on the stable isotopic signatures of different waters sources and the plant stem water (Dawson and Ehleringer, 1991; Busch et al., 1992; Alstad et al., 1999; Zhou et al., 2017; White and Smith, 2020). Nevertheless, For example, riparian *Liquidambar styraciflua* growing along a perennial stream took up river water by 30–35% in the southern Appalachian foothills, USA (White and Smith, 2020). Alstad et al. (1999) found that riparian *Salix* relied on rivers for approximately 80% of its water, which made it vulnerable to changes in river water and hydrological conditions on the northeast side of Rocky Mountain National Park, Colo.-

However, there was a debate on whether the river water is a potential water source for riparian trees or not and 70 how it became available to plants. Ssome studies argued that the river water was not a direct potential water source and rarely contributed to riparian trees (Dawson and Ehleringer, 1991; Bowling et al., 2017; Dawson and Ehleringer, 1991; Wang et al., 2019a). Dawson and Ehleringer (1991) firstly discovered that the mature streamside trees growing in or next to a perennial river did not use river water but depended on the water sources from deeper strata. This Similar finding has also been foundproven in riparian phreatophytic trees (Populus fremontii and Salix 75 gooddingii) and riparian deep-rooted tree species (Busch et al., 1992; Bowling et al., 2017; Wang et al., 2019a). Even under shallow groundwater with high salinity, no river water was directly absorbed taken up by riparian Eucalyptus coolabah alongside an ephemeral arid zone river in Australia (Costelloe et al., 2008). Growing evidence showed that riparian trees rarely took up river water directly at a certain distance away from the riverbank because their lateral roots could not reach the river (Mensforth et al., 1994; Thorburn and Walker, 1994); 80 Nevertheless, riparian trees could indirectly utilize the river water that recharges riparian deep zone (e.g., deep soil water and groundwater) when their roots tap into the groundwater level (Mensforth et al., 1994; Thorburn and Walker, 1994; Wang et al., 2019a). If we take the-river water as a direct water source, the RWC to transpiration flux of riparian trees may be overestimated. HOther studies claimed that river water merging into deep riparian

- 85 (Mensforth et al., 1994; Wang et al., 2019b). It remained unclear that how much river water exactly contributed to riparian trees nearby a losing river. This might lead to inaccurate estimations when river water could not be directly accessed by lateral roots of riparian trees growing at a certain distance away from the riverbank (Mensforth et al., 1994; Thorburn and Walker, 1994). How to separate and quantify the indirect contributions of the indirect river water source to riparian trees nearby losing rivers is a great challenge.
- 90 The graphical inference and direct comparison of stable isotopic values between plant stem water and different water sources (Dawson and Ehleringer, 1991; Busch et al., 1992; Costelloe et al., 2008; Zhao et al., 2016), statistical two- or multi-source linear mixing models (Alstad et al., 1999; Zhou et al., 2017), and the MixSIAR Bayesian mixing model (Wang et al., 2019a; Wang et al., 2020; White and Smith, 2020; Li et al., 2021) integrated with-the stable water-isotopes ( $\delta^{2}$ H and  $\delta^{18}$ O) have been widely used to identify the potential water sources taken
- 95 up by riparian trees. The MixSIAR model has more advantages in quantifying water source contributions and accounting for uncertainties in the isotopic values (Stock and Semmens, 2013; Ma et al., 2016). The RWC to riparian trees can be estimated indirectly by quantifying both the direct water source contributions to riparian trees and the RWC to riparian deep water. A multi-iteration method (Marek et al., 1990; Zaid, 2010) is key to calculate the proportional contributions of total (old and current) river water to riparian deep water, which improves the
- 100 estimation accuracy of the RWC to riparian trees. The radioactive isotope (<sup>222</sup>Rn) has been widely used for tracing groundwater origins and corresponding pathways in the riparian zone (Close et al., 2014; Zhao et al., 2018). It is helpful to estimate the residence time of recharged groundwater from river water and its effects on the RWC to riparian trees. A combination of these methods can give a more reliable quantification of the RWC to riparian trees.\_
- 105 The RWC to riparian trees could substantially affect the leaf-level water use efficiency (WUE) and healthy growth of riparian trees. The tTree WUE is a key characteristic of plant water use, which can be defined as the ratio of photosynthetic rate to transpiration rate. Since leaf δ<sup>13</sup>C values are positively related to tree WUE, the-leaf δ<sup>13</sup>C has been widely used as an indicator of tree WUE for C<sub>3</sub> photosynthesis plants (Farquhar et al., 1989). For example, Thorburn and Walker (1994) found that the riparian *Eucalyptus camaldulensis* beside the ephemeral stream had higher tree WUE with more frequent access to river water based on the leaf δ<sup>13</sup>C measurements. Moreover, the fluctuation of the depth to the water tablewater table depth (WTD) in the riparian zone resulting

Clark, 2001; Liu et al., 2017; Xia et al., 2018). However, little attention has been paid to quantifying the relationships between the RWC to riparian trees and the tree WUE <u>as well as</u>or WTD near a losing river.

- 115 The river water contribution (RWC) to riparian trees has been widely estimated using the data comparison, graphical inference, two or multi source linear mixing models and Bayesian mixing models (MixSIR, SIAR, SISUS, MixSIAR) accompanied with stable water isotopes ( $\delta^2 H$  and  $\delta^{18}O$ ) (Dawson and Ehleringer, 1991; Ehleringer and Dawson, 1992; White and Smith, 2020). Several previous studies reported that the WUE of riparian trees varied with the flucuation of water table depth (WTD) in the riparian zone due to changing river flow (Horton and Clark, 2001; Liu et al., 2017; Xia et al., 2018).
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The RWC could substantially affect the leaf level water use efficiency (WUE) and healthy growth of riparian trees. The WUE is a key characteristic of plant water use, which can be defined as the ratio of photosynthetic rate to transpiration rate. Since leaf  $\delta^{13}$ C values are positively related to WUE, the leaf  $\delta^{13}$ C has been widely used as an indicator of WUE (Farguhar et al., 1989). Thorburn and Walker (1994) found that the riparian Eucalyptus 125 camaldulensis beside the ephemeral stream had higher WUE with more frequent access to river water based on the leaf  $\delta^{13}$ C measurements. Nevertheless, Sun et al. (2008) observed that the river water availability had little effect on WUE because there was no significant difference in WUE between riparian Pinus massoniana and nonriparian Pinus quercus. Several previous studies reported that the WUE of riparian trees varied with the flueuation of water table depth (WTD) in the riparian zone due to changing river flow (Horton and Clark, 2001; Liu et al., 130 2017; Xia et al., 2018).-For example, Horton and Clark (2001) showed that the WUE of riparian Salix gooddingii and Populus fremontii increased with increasing WTD, and riparian Tamarix chinensis had significantly higher WUE under fluctuated deep WTD in dry year than those under constant shallow WTD in wet year. In comparison, the WUE of riparian species decreased significantly along a gradient of increasing WTD (from 0.5 m to 12 m) in the middle reaches of Heihe River Basin, China (Liu et al., 2017). However, little attention has been paid to 135 quantifying the relationships between the RWCs to riparian trees and WUE as well as WTD.

The overall aim of this study was to clarify the effects of river water on water use of riparian trees along a gradient of WTD. Focusing on a losing river in Beijing, China, the specific objectives of this study were to: (1) to propose an iteration method together with the MixSIAR model and water stable isotopes ( $\delta^{2}$ H and  $\delta^{18}$ O) to quantify the RWCs; (2) to-determine the proportional contributions of river water to riparian trees at different 140 distances away from the riverbank; (3) to-identify the relationships between the RWCs to riparian trees and WUE (indicated by leaf  $\delta^{13}$ C values) as well as WTD. These results will provide critical insights into plantation management, bank storage conservation and ecosystem and healthy ecosystem enhancement for losing rivers.

#### 2 Materials and methods

#### 2.1 Study area

- The study area was in the reaches of the Chaobai River, located in Shunyi district, Beijing, China (40°07'30"N, 116°40'37"E) (Fig. 1). The temperate continental sub-humid monsoon climate prevails in this area, with an annual mean temperature and evaporation of 11.5 °C and 1175 mm, respectively. <u>The average total precipitation from</u> <u>April to November between 1961 and 2021 is 532.8 mm, with 84.5% of which falling in rainy season (from June</u> <u>to September) (Fig. 2a).</u> Due to continuous drought and groundwater overexploitation, the Chaobai River dried
- up during from 1999 to 2007 and the riparian ecosystem seriously degraded seriously. The "ecological water" (including reclaimed water, reservoir water, and diverted water by the South-to-North Water Transfer Project) has been supplied via a systematic water release by dams to restore this dry river since 2007.5 A total of 51.1 million and 380 million cubic meters of ecological water sources were released to the Chaobai River in 2019 and 2021, respectively. Mand more than 33 km<sup>2</sup> of the riparian zone has been reeovegetatred with different tree species by until 2020. The *Salix S. babylonica* (L.) was one of the most widely planted species alongside the Chaobai River. Three plots at distances of 5 m (D05), 20 m (D20), and 45 m (D45) away from the riverbank were selected for field measurements and sample collection (Fig. 1).

#### 2.2 Field measurements and data collection

The field measurements were conducted <u>during\_from</u> April to November in 2019 and 2021, with no field observation in 2020 due to the COVID-19. The daily precipitation data from 1961 to 2021 <u>and the daily mean</u> temperature (T), relative air humidity (RH), solar radiation and reference evapotranspiration (ET<sub>0</sub>) data during the <u>observation period</u> in the Shunyi district <u>was-were</u> collected from the China Meteorological Data Service Centre (http://data.cma.cn/en). <u>Daily mean vapour pressure deficit (VPD) was calculated using the RH and T data</u> (Schoppach et al., 2019).

# 165The groundwater levels in each plot were recorded monthly in 2019 and 2021 via the a pressure stage gauge(HOH-S-Y, King Water Co Ltd., Beijing, China) installed in the groundwater monitoring well. The river water

level was recorded using a water level gauge at the same time with the observation of groundwater levels.

The average total precipitation during April to November between 1961 and 2021 is 532.8 mm (Fig. 2a). The observation period in 2021 was wet with total precipitation of 802.5 mm, which was 1.8 times of that in dry year of 2019 (445.6 mm) (Fig. 2b and c). The river water level fluctuated at 27.9-28.9 m in 2019 and 27.3-29.7 m in 2021 (Fig. 3). The mean WTD in three plots in 2019 (2.7 ± 0.3 m) was significantly larger than that in 2021 (1.7  $\pm$  0.5 m) (p < 0.05). The WTD decreased with increasing distances from the riverbank in both 2019 and 2021.

#### 2.3 Sample collection and isotopic analyses

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Twelve sampling campaigns on May 5, June 14, July 26, August 15, September 26, November 5 in 2019 and April 175 24, May 25, June 26, July 15, September 1, November 5 in 2021 were conducted to collect groundwater, precipitation, river water, soil, stem, and leaf samples. Groundwater in each plot was sampled by a sucking pump from the monitoring well, and a plexiglass hydrophore water sample collector with capacity of 1L was used to collect the nearby river water. Precipitation was sampled after each precipitation event via a device consisting of a funnel, a polyethylene bottle, and a ping-pong ball. A total of 135 precipitation samples were collected 180 throughout the whole years of 2019 and 2021 in the observation period via a device consisting of a funnel, a polyethylene bottle and a ping pong ball. All precipitation, groundwater, and river water samples were stored in a refrigeratingon box with several ice bags to avoid minimize evaporation effects in the field, then they were delivered to the laboratory and kept at 4 °C in the refrigerator until water isotope ( $\delta^2$ H and  $\delta^{18}$ O) analysis. The groundwater and river water were also collected with the-100-ml brown glass vials bottles to measure the-222Rn 185

concentration in the study areafield.

Three-One riparian S. babylonica trees was chosen (with mean diameter of  $28.6 \pm 4.4$  cm at breast height) growing in three each plots were chosen on each sampling campaign for  $\delta^2 H$  and  $\delta^{18} O$  measurements in xylemstem water as well as  $\delta^{13}$ C analysis in plant leaves. The mean diameter at breast height of three sampled trees was 28.6  $\pm$  4.4 cm. Five mature and Several suberized stem samples were taken firstly cut from the same riparian S. babylonica tree in each plot using an averruncator with the length of 5 m. We<sub>7</sub> removed the bark and phloem of the sampled stems, and then put the remaining xylem samples into a 12-ml brown glass vial sealed with the parafilm. Meanwhile, the mature leaves without petioles were sampled from the collected stems using pruning shears.stored at -10 °C until water isotope analysis. The remaining xylem and mature leaf samples were stored in a refrigerationing box with several ice bags in the field. Then the xylem samples were transported to the laboratory

195 and kept in a refrigerator at -10 °C before water extraction and isotope analysis. The mature leaves were sampled from the collected stems, oven-dried at 65 °C for 72 h on the day of sampling, then grinned grinded and passed through a 0.15 mm sieve to analyze leaf  $\delta^{13}$ C (Wang et al., 2019b; Cao et al., 2020; Wang et al., 2019b).

Soils at depths of 0–5, 5–10, 10–20, 20–30, 40–60, 60–80, 90–110, 150–170, 190–210, 250–270, and 280–300 cm in one soil profile nearby the selected *S. babylonica* trees were sampled by a power auger (CHPD78,

200 Christie Engineering Company, Sydney, Australia). One <u>part-portion</u> of each soil sample was put into a 12-ml brown glass vial and stored at −10 °C for before water stable isotope analysis, and the other <u>part-portion</u> was packed into an aluminum box for gravimetric soil water content (SWC) measurement via the oven-drying method (Wang et al., 2019b; Li et al., 2021; Wang et al., 2019b).

The automatic cryogenic vacuum distillation system (LI-2100, LICA, Beijing, China) was used to extract water in-from xylstem and soil samples, which generally ran for at least 2.5 h. We weighted all the xylem and soil samples before and after extraction. Subsequently, the efficiency of water extraction was calculated in order to ensure the water extraction efficiency above 99% and noto avoid isotopic fractionation during the-water extraction process and kept the efficiency of water extraction more than 99% to ensure no isotopic fractionation. The δ<sup>2</sup>H and δ<sup>18</sup>O in soil water, river water, groundwater, and precipitation were analyzed through an isotopic ratio infrared

spectroscopy system (IRIS) (DLT-100, Los Gatos Research, Mountain View, USA) (Li et al., 2021). The Isotope Ratio Mass Spectrometry system (IRMS) (MAT253, Thermo Fisher Scientific, Bremen, Germany) which could prevent from organic pollution of plants was used to measure δ<sup>2</sup>H and δ<sup>18</sup>O in stem-xylem water as well as leaf δ<sup>13</sup>C value. There was the same measurement accuracy of ±1‰ for δ<sup>2</sup>H and ± 0.1‰ for δ<sup>18</sup>O between for both the IRIS and IRMS systems (±1‰ for δ<sup>2</sup>H and ± 0.1‰ for δ<sup>18</sup>O). The Vienna Standard Mean Ocean Water (VSMOW) was used to calibrate and normalize the δ<sup>2</sup>H and δ<sup>18</sup>O measurements in different waters, while the Vienna Pee Dee Belemnite (V-PDB) was used for calibrating leaf δ<sup>13</sup>C values.

The <sup>222</sup>Rn concentration in the groundwater and river water samples (C<sub>Water</sub>, Bq/l) was determined based on the <u>air</u> <sup>222</sup>Rn concentration <u>values</u> (C<sub>Air</sub>, Bq/m<sup>3</sup>) measured by a <sup>222</sup>Rn monitor (Alpha GUARD PQ2000 PRO, Bertin Instruments, Germany) (C<sub>Air</sub>, Bq/m<sup>3</sup>). 100 ml of the water sample was slowly poured into the air-tight glass bottles and then purged with air in a closed gas <u>cycle\_cycling\_system</u>. The\_\_C<sub>Air</sub>\_<u>-in the</u> <sup>222</sup>Rn monitor was recorded at <u>a</u> 10-minute interval<del>s</del>, The air inside the measurement set-up had commaintained a certain <sup>222</sup>Rn concentration right before <u>injecting</u> the water sample injection (C<sub>System</sub>, Bq/m<sup>3</sup>). It is generally assumed that the already existing C<sub>System</sub> can be ignoredignored accordingly when C<sub>System</sub> is around or lower than 80 Bq/m<sup>3</sup>. In this study, and more than four intervals were conducted to ensure that the C<sub>System</sub><sup>222</sup>Rn concentration in the measuring
 set up before sampling (background) (C<sub>System</sub>, Bq/m<sup>3</sup>) was less than 80 Bq/m<sup>3</sup>.–\_The measurement range of C<sub>Air</sub> was 2–2,000,000 Bq/m<sup>3</sup> with a measurement precision of 3%. The above measured C<sub>Air</sub> value was not yet the <sup>222</sup>Rn concentration in the measured water sample (C<sub>water</sub>), because the <sup>222</sup>Rn driven out had been diluted by the air within the <sup>222</sup>Rn monitor and a small part ortion of the <sup>222</sup>Rn remained diluted in the watery phase. The C<sub>water</sub> c C<sub>water</sub> can be calculated as:\_

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$$C_{\text{Water}} = \frac{C_{\text{Air}} \times \left(\frac{V_{\text{System}} - V_{\text{Sample}}}{V_{\text{Sample}}} + k\right) - C_{\text{System}}}{1000}$$
(1)

where  $V_{\text{System}}$  is the interior volume of the measuring set-up (ml), which is 1122 ml in this study.  $V_{\text{Sample}}$  is the volume of water sample (ml). k is the <sup>222</sup>Rn distribution coefficient of water/air (–), which can be set as 0.26 within the specified temperature range around a mean room temperature of 20 °C (Clever, 1985).

<u>In this study, the average residence time (T<sub>res</sub>, day) of recharged groundwater recharged from the river water</u>
 <u>to the underlying aquifer and/or riverbank-was also identified based on the <sup>222</sup>Rn isotopes (Hoehn and Von Gunten, 1989), which was described as follows:</u>

$$T_{\rm res} = \frac{1}{\lambda} \times \ln\left(\frac{C_{\rm e} - C_{\rm r}}{C_{\rm e} - C_{\rm g}}\right)$$
(2)

where λ represents the decay coefficient (0.181\_day<sup>-1</sup>) (Hoehn and Von Gunten, 1989). C<sub>e</sub> represents the <sup>222</sup>Rn concentration of background groundwater when the equilibrium between radon production and decay is reached.
 (7400 Bq/m<sup>3</sup> in this study): The measuring <sup>222</sup>Rn concentration of groundwater in aquifers more than 100 m away from the riverbank remains constant in this study (with an average value of 7400.0 ± 35.4 Bq/m<sup>3</sup>), suggesting that C<sub>e</sub> can be defined as 7400.0 Bq/m<sup>3</sup>. C<sub>r</sub> represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>); C<sub>g</sub> represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>); C<sub>g</sub> represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>); C<sub>g</sub> represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>); C<sub>g</sub> represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>); C<sub>g</sub> represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>); C<sub>g</sub> represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>); C<sub>g</sub> represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>); C<sub>g</sub> represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>); C<sub>g</sub> represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>); C<sub>g</sub> represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>); C<sub>g</sub> represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>).

#### 2.4 Determination of RWC to riparian trees

245 Riparian trees at a certain distance away from the riverbank rarely used river water directly, as their lateral roots could not reach the river (Mensforth et al., 1994; Thorburn and Walker, 1994). Nevertheless, riparian trees could continuously absorb river water that merged into riparian deep water (including groundwater and deep soil water within the capillary fringe) when their deep roots tapped into the water table. In this study, stable water stable isotopes (δ<sup>2</sup>H and δ<sup>18</sup>O) integrated with the MixSIAR model were integrated within the MixSIAR model and an

250 <u>iteration method and an iteration method wereas</u> proposed to identify the original (before and during the observation period)-contributions of <u>the indirect</u> river water that <u>merged-rechargedinto</u> riparian deep water to riparian *S. babylonica* trees (Figs. 4 and 5). Firstly, the <u>direct water source (including soil water in different layers and groundwater) contributions to riparian trees root water uptake patterns</u> were determined via δ<sup>2</sup>H and δ<sup>18</sup>O in different waters and the MixSIAR model, <u>without considering river water as a direct water source for riparian trees</u>. Secondly, the proportional contributions of river water to riparian deep water <u>(i.e., riparian groundwater and deep soil water in the 80–170 cm layer</u>) were figured determined out by the MixSIAR model and water isotopes. Finally, the proposed iteration method was used to quantify the proportions of the <u>indirect river water source taken</u>.

<u>up by RWC to riparian trees (Figs. 4 and 5)</u>.

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The MixSIAR model is a Bayesian mixing model which can be integrated with the stable isotopes to quantify
 the proportions of source contributions to a mixture (Stock and Semmens, 2013). The input data of the MixSIAR model include mixture data, source data, and discrimination data. In this study, the mean and standard deviation (SD) of the isotopic values of each water source for riparian trees/riparian deep water were input as source data into the MixSIAR, while the measured isotopic values of xylem water/riparian deep water were input as raw mixture data into the MixSIAR. The discrimination data for both δ<sup>2</sup>H and δ<sup>18</sup>O were set to zero, because the input δ<sup>2</sup>H and δ<sup>18</sup>O values in the MixSIAR were non-fractionated or δ<sup>2</sup>H-corrected. The Markov Chain Monte Carlo

parameter was set to the run length "very long". Both the trace plots and three diagnostic tests (i.e., Gelman– Rubin, Heidelberger–Welch, and Geweke) were used to determine whether the MixSIAR model converged or not (Stock and Semmens, 2013). Then, the mean and SD values of different water source contributions <u>-(median</u> <u>values)</u> could be estimated with the MixSIAR model.

### 270 **2.4.1** Quantifying <u>proportional direct water source</u> contributions <u>of direct water sources</u> to riparian trees

In this study,<u>S</u>-soil water at different depths, which was mixed proportionally with precipitation, old soil water, or even river water and groundwater, was taken up by riparian *S. babylonica* directly. We measured soil water isotopes at 11 depths in the three plots. In order to reduce errors in the analytical procedure, four soil layers (0–30 cm, 30–80 cm, 80–170 cm, and 170–300 cm) were dividused to identify the main root water uptake depth of riparian trees according to seasonal variations in the SWC, water isotopes and WTD. The average soil water

	isotopes values at depths of 0-5 cm, 5-10 cm, 10-20 cm, and 20-30 cm were calculated for the 0-30 cm soil
	layer, because the water isotopes went through strong evaporation and the-SWC varied significantly seasonally.
	The soil water isotope values at depths of 40-60 cm and 60-80 cm were averaged for the 30-80 cm soil layer, and
280	those values at 90-110 cm and 150-170 cm depths were averaged for the 80-170 cm soil layer assince the water
	isotopes and SWC were relatively stable. The average isotopic values of soil water at deep depths (190-210 cm,
	250-270 cm, and 280-300 cm) were calculated for the 170-300 cm soil layer, which varied with the fluctuations
	of the groundwater levels. Groundwater also could -be regarded as a relatively direct stable water source for
	phreatophyte riparian trees (Dawson and Ehleringer, 1991; Busch et al., 1992). In terms of seasonal variations in
285	the SWC, water isotopes and WTD, four soil layers (0-30, 30-80, 80-170, and 170-300 cm) were divided to
	identify the main root water uptake depth of riparian trees in the three plots (Figs. 2, 3 and S1). As the isotopic
	composition of soil water in the 170-300 cm layer was similar to that of groundwater, they were considered to be
	one water source (groundwater). Mensforth et al. (1994) and Thorburn and Walker (1994) characterized the outer
	projected edge of canopy as the extension range of lateral roots, which could indicate whether riparian trees take
290	up river water directly or not. In this study, the outer-projected edge of canopy was less than 5 m for riparian S.
	babylonica tree closest to the river (5 m away from the riverbank). It indicated that the lateral roots of S. babylonica
	trees could not tap into the river. Therefore, the river water was not considered as a direct potential water source
	for tree water uptake, while groundwater and soil water in the 0-30, 30-80, and 80-170 cm layers, and
	groundwater-were determined used as the direct potential water sources for riparian S. babylonica.
295	In this study, the $\delta^2$ H offsets between the xylem water in riparian trees and its corresponding potential source
	waters were observed, which could be resulted by the from $\delta^2$ H fractionation in the plant water use processes (Li
	et al., 2021; Cernusak et al., 2022). These $\delta^2$ H offsets could lead to biglarge errors in estimating the water source
	<u>contributions using the MixSAIR model. IDue n order to eliminate the <math>\delta^2</math>H offsets of stem xylem water from its</u>
	potential <u>water</u> sources, the measured <u>stem-xylem</u> water $\delta^2 H$ values were corrected via the potential water source
300	line (PWL) proposed by Li et al. (2021). The PW-excess (PW-excess = $\delta^2 H - a_p \delta^{18} O - b_p$ ; $a_p$ and $b_p$ were the slope
	and the intercept of the PWL, respectively) was calculated to indicate the $\delta^2 H$ deviation from the PWL, which
	was subsequently subtracted from the measured xylem water $\delta^2 H$ values. The corrected $\delta^2 H$ and raw $\delta^{18}O$ and
	corrected $\delta^2$ H in stem <u>xylem</u> water were set as the mixture data in the MixSIAR model to quantify the direct water
	source contributions of direct water sources to riparian S. babylonica. The parameter settings of the MixSIAR
I	

305 model have been described in detail by Stock and Semmens (2013) and Li et al. (2021). The contributions of riparian deep soil water in 80–170 cm layer to riparian trees (P<sub>s</sub>) and the groundwater contributions to riparian trees (P<sub>s</sub>) could be determined in particular to evaluate the RWCs to riparian trees.

## 2.4.2 Identifying <u>Quantifying</u> water source <u>contribution</u>s for to deep soil water and groundwater

- 310 Riparian deep soil water and groundwater could be continuously recharged by river water when the groundwater levels lied below the riverbeds (i.e., losing rivers). The MixSIAR model in conjunction with water isotopes ( $\delta^2$ H and  $\delta^{18}$ O) were <u>applied</u>-used to quantify the proportional contributions of current (between previous sampling time t-1 and current sampling time t) river water to riparian deep water (i.e., deep soil water in the 80-170 cm layer or groundwater). RWCs to riparian deep water. As shown in Fig. S2a, the The potential water sources of 315 riparian deep soil water in the 80–170 cm layer at <del>current sampling time (t)</del> included the in-situ (i.e., water that is already in the deep soil layer or groundwater) -soil water in this layer at previous sampling time (t-1), soil water in the 0-80 cm layer at t-1, river water between t-1 and t, precipitation between t-1 and t, and groundwater between t-1 and t\_(Fig. 4a). The potential water sources for riparian groundwater at t were considered as the in-situ groundwater at t-1, soil water in the 0-170 cm layer at t-1, river water between t-1 and t, and precipitation between 320 t-1 and t (Fig. <u>S2b4b</u>). The  $\delta^{2}$ H and  $\delta^{18}$ O values in riparian deep water (deep soil water in 80–170 cm layer or groundwater) at t were set as the mixture data in the MixSIAR model, while the water isotopes of their potential water sources were considered as the source data.

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In this study, the average residence time (T<sub>res</sub>) of groundwater recharged from the river to the underlying aquifer and/or riverbank was also identified based on the <sup>222</sup>Rn isotopes, which was described as follows:

$T = \frac{1}{2} \times \ln \left( \frac{C_e - C_f}{C_f} \right)$		
$\frac{1}{\text{res}} = \frac{1}{\lambda} \frac{1}{$	(2)	

where  $\lambda$  represents the decay coefficient (0.181 d<sup>±</sup>).  $C_{e}$  represents the <sup>222</sup>Rn concentration when the equilibrium between radon production and decay is reached (7400 Bq/m<sup>3</sup> in this study);  $C_{e}$  represents the <sup>222</sup>Rn concentration of river water (Bq/m<sup>3</sup>);  $C_{e}$  represents the <sup>222</sup>Rn concentration of groundwater (Bq/m<sup>3</sup>).

#### **330 2.4.3 An iteration method to determine RWCs to riparian trees**

The proportional contributions of the river water between t-1 and t to riparian trees could be quantified when

riparian deep-water contributions to trees and the RWCs to riparian deep water were both figured outdetermined. It was worth notinged that <u>Rriparian deep soil water (80-170 cm)</u> and groundwater couldan be continuously recharged by river water continuously when the groundwater levels lied below the riverbeds (i.e., losing rivers).

Therefore, the proportional contribution of the old river water (before t-1) to riparian deep water should not be ignored. The total RWC to riparian deep water should be quantified explicitly during the entire period of the river losing flow to riparian deep zone since 2007. the riparian in situ deep water (i.e., in situ 80–170 cm soil water and in-situ groundwater) were also merged by old river water before t-1. In this study, We assumed that the contributions of old river water before t-1 to riparian in-situ deep water were assumed to be identical to consistent with those proportions contributions of current river water (between t-1 and t) to riparian in-situ deep water (i.e., "s<sub>r</sub>" and "g<sub>r</sub>"). An iteration method was proposed to quantify the original total RWC to riparian *S. babylonica* trees nearby the losing rivers, which was described as follows:

$$RWC = P_s * S_r + P_g * G$$

(3)

$$= P_s^* (s_r^t + s_r^{t-1} s_r + s_{\overline{r}}^{\theta}) + P_g^* (g_r^t + g_r^{t-1} g_r + g_{\overline{r}}^{\theta})$$

$$345 = P_{s}^{*}(s_{r}^{t} \pm s_{r}^{t+s}s_{s}^{t-1} \pm s_{r}^{t}(s_{s}^{t-1})^{2} \pm s_{g}^{t} \pm s_{g}^{t} \pm s_{g}^{t} \pm s_{g}^{t} \pm s_{g}^{t} \pm s_{g}^{t+s}(s_{g}^{t-1})^{2} + s_{r}^{t} \pm s_{s}^{t} \pm s_{g}^{t} \pm s_{g}^{t} \pm s_{g}^{t} \pm s_{s}^{t} \pm$$

355

where S<sub>r</sub> and G<sub>r</sub> represent original total (before and during the observation period) RWCs to riparian deep soil water in the 80–170 cm layer and groundwater, respectively. The P<sub>s</sub> and P<sub>g</sub> represent the contributions of riparian deep soil water in the 80–170 cm layer and groundwater to riparian trees, respectively. The s<sub>r</sub><sup>t-1</sup> and g<sub>r</sub><sup>t-1</sup> and g<sub>r</sub><sup>d-1</sup> an 

#### 2.5 DataStatistical analysis

One-way analysis of variance (ANONA) incorporating with the Kolmogorov-Smirnov, Levene's and post-hoc
Tukey's tests (*p* < 0.05) were used to investigate the statistic differences of different variables. The variables included The statistic differences in the WTDs, SWC, δ<sup>2</sup>H and δ<sup>18</sup>O in different water-bodie sources and xylems,
<sup>222</sup>Rn concentration of river water and groundwater, contributions of different potential-water sources to riparian deep water or trees-contributions, and leaf δ<sup>13</sup>C values among in the three plots in 2019 and 2021-were analyzed by. The linear regression model was fitted to the whole dataset in both two years to get the general relationships
(not only available for the observation periods) between the WTD, leaf δ<sup>13</sup>C values and post hoc Tukey's tests (*p* < 0.05). The relationships between the WTD, leaf δ<sup>13</sup>C values and RWCs to riparian trees were determined by the regression analysis method. The statistical analysis was performed in the Excel (v2016) as well asnd SPSS (24.0, Inc., Chicago, IL, USA).

#### 380 **3 Results**

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#### 3.1 Hydro-meteorological conditions

The observation period in 2021 was wet with a total precipitation of 802.5 mm, which was 1.8 times higher than for the drier year 2019 (445.6 mm) (Fig. 2a). The precipitation amount during rainy season accounted for 75.4% and 97.0% in 2019 and 2021, respectively. The annual mean temperature during the observation period in 2019 and 2021 was 22.4 °C and 21.8 °C, respectively. The daily mean VPD and  $ET_0$  increased during the observation period, reaching a peak in June and May, respectively (Fig. S1). The average daily VPD during the observation period was significantly higher in dry 2019 (1.1 KPa) than in wet 2021 (0.9 KPa) (p < 0.05) (Fig. S1a and b). There was a significant difference in the average daily ET<sub>0</sub> from June to September between dry 2019 (5.0 mm/day) and wet 2021 (4.3 mm/day) (p < 0.05), but no significant difference was observed during the remaining observation period between the two years (p > 0.05) (Fig. S1c and d). No significant difference in the daily mean net radiation during the observation period was found between dry 2019 and wet 2021 (p > 0.05) (Fig. S1 c and

<u>d).</u>

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The river water level fluctuated between 27.9 m and 28.9 m in 2019 and between 27.7 m and 29.3 m in 2021 (Fig. 3). The mean WTD across the three plots was significantly (p < 0.05) highdeeper in 2019 ( $2.7 \pm 0.3$  m) than in 2021 ( $1.7 \pm 0.5$  m). The WTD increased with increasing distances from the riverbank in both 2019 and 2021 (Fig. 3). The river water continuously recharged the groundwater system ("losing" river) during the observation periods in 2019 and 2021, indicated by a lower groundwater level than the river water level (Fig. 3). Significantly higher SWC was observed in 2021 compared to 2019 (p < 0.05) (Fig. S2). Significantly higher SWC was observed in 2021 compared with that to in 2019 (p < 0.05) (Fig. S12). The SWC of of each all four soil layers at D45 was significantly lower than that at D05 and D20 in 2021 (p < 0.05), while no pronounced difference in the SWC of in the 0–30 cm layer was observed among the three plots in 2019 (p > 0.05) (Fig. S42).

#### 3.1-2 Direct water source contributions to Water uptake patterns of riparian trees

It was evident in Fig. 5 that the δ<sup>2</sup>H and δ<sup>48</sup>O in precipitation Precipitation was significantly more depleted in δ<sup>2</sup>H and δ<sup>18</sup>O in 2021 (-52.9‰ ± 30.2‰ for δ<sup>2</sup>H and -8.1‰ ± 3.8‰ for δ<sup>18</sup>O) than those in 2019 (-29.2‰ ± 18.8‰ for δ<sup>18</sup>O) for δ<sup>2</sup>H and -4.1‰ ± 3.0‰ for δ<sup>18</sup>O) (p < 0.05) (Fig. 6). The significantly larger slope of the Local local Meteorie meteoric Water water Line-line in 2021 (7.8) was significantly higher with respect than to that in 2019 (5.5) (p < 0.05), which also suggested that the falling raindrops undergone stronger sub-cloud evaporation in 2019 (Zhao et al., 2019) more depleted precipitation isotopes in 2021 (p < 0.05). Significantly higher SWC was observed in 2021 compared with that in 2019 (p < 0.05) (Fig. S1). The SWC of all four soil layers at D45 was significantly lower than that at D05 and D20 in 2021 (p < 0.05) (Fig. S1). The SWC of all four soil layers at D45 was significantly lower was observed among three plots in 2019 (p > 0.05) (Fig. S1). The δ<sup>2</sup>H and δ<sup>18</sup>O in the superficial surface soil water in (above θ=30 cm layer depth) in 2021 were significantly more depleted lower and more variable in 2021 -than those in 2019 (p < 0.05) (Fig. 56). NeverthelessIn contrast, there were slightly enriched-higher water isotopes-isotopic</li>

compositions in the 30-170 cm soil layer in 2021 compared with those to in 2019. No significant difference in the

415 isotopic compositions of <u>the</u> soil water below 170 cm depth and groundwater <u>waswere</u> observed between 2019 and 2021 (p > 0.05). <u>The δ<sup>2</sup>H and δ<sup>18</sup>O in soil water in the 80–170 cm layer were significantly lower than groundwater in 2019 (p < 0.05)</u>, while no significant difference was observed between soil water isotopes in the <u>80–170 cm layer and groundwater isotopes in 2021 (p > 0.05)</u>. As shown in Fig. 5, the isotopes in <u>gG</u>roundwater were was significantly more depleted in δ<sup>2</sup>H and δ<sup>18</sup>O than compared to those in river water in both two years (p < 0.05) (Fig. 6). The δ<sup>2</sup>H and δ<sup>18</sup>O in stem-xylem water during the observation periods in 2019 and 2021 were not significantly different (p > 0.05), but they were gradually <u>depleted lower</u> with the increasing distances from the riverbankin riparian zone.

The contributions of the superficial surface soil water (above 30 cm depth) to transpiration flux of riparian trees in 2019 (20.1% ± 9.7%) were similar to those in 2021 (19.0% ± 10.5%). As shown in Fig. 6, n No significant difference in the soil water contributions to riparian *S. babylonica* was also observed in the 30–80 cm layer between the two years (p>0.05) As shown in (Fig. 67). The *S. babylonica* tree species principally relied on riparian deep water below the 80 cm depth in both 2019 (55.9%) and 2021 (57.1%). There was no significant difference in the riparian deep-water contributions to the transpiration flux of *S. babylonica* trees between the three distances from the riverbank (p > 0.05) (Fig. 7). Nevertheless, tThe soil water contributions in the 80–170 cm layer to riparian trees reduced decreased with the increasing distance away from the riverbank in both years, whereas the proportions of groundwater contributions taken up by riparian trees increased from D05 to D45 in both 2019 (from 27.6% to 32.1%) and 2021 (from 17.0% to 32.2%) (Fig. 67). It was found that the groundwater contributions to riparian *S. babylonica* trees increased evidently significantly (p < 0.05) from April to July; Nevertheless, but tThey plummeted significantly (p < 0.05) and reached minimum in September in 2021.

#### 435 **3.2-3** Water source contributions to riparian deep soil water and groundwater

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Significant seasonal and interannual variations of different water source contributions to riparian deep soil water in 80–170 cm layer were found during the observation periods in the study area (p < 0.05). RThe primary water sources of riparian deep soil water in the 80-170 cm layer were as primarily recharged by the in-situ soil water in this layer (mean of 33.1%) and groundwater capillary rise (mean of 25.3%) in 2019 (Fig. 78). In comparison, the in- situ soil water in the 80-170 cm layer (mean of 23.9%), groundwater capillary rise (mean of 24.6%), and river water (mean of 24.4%) contributed evenly to riparian deep soil water in 2021. The in–situ soil water contribution to riparian deep soil water in 2019 was significantly higher in 2019 than that in 2021 (p < 0.05). However, the river water contributed less to riparian deep soil water in 2019 (mean of 15.7%) compared with to 2021 (p < 0.05). The RWC to riparian deep soil water was the lowest in August in 2019 ( $11.3\% \pm 4.5\%$ ) but and in June in 2021 ( $13.6 \pm 3.8\%$ ), respectively. The in--situ soil water contributions to riparian deep soil water showed a significant increase with increasing distance away from the riverbank, while the RWCs to riparian deep soil water decreased from D05 to D45 in both years (p < 0.05) (Fig. 78).

There were significant differences in interannual and seasonal water source contributions to riparian groundwater in three plots in dry 2019 and wet 2021 (p < 0.05). The in-situ groundwater was the main source of 450 riparian groundwater in both years (Fig. 89), contribution but its contribution was significantly higher in 2019 (mean of  $56.0\% \pm 11.2\%$ ) than that in 2021 ( $37.1\% \pm 16.7\%$ ) (p < 0.05) (Fig. 9). The average contribution of the river water to riparian groundwater was recharged by river water with mean of  $28.1\frac{6}{2} \pm 12.1\%$  during the observation period. There was a significantly higher RWC to riparian groundwater in 2021 (mean of  $35.1\frac{10}{2}$  ± 11.9%) than that in 2019 (mean of  $21.1\% \pm 7.2\%$ ) (p < 0.05). The lowest RWC ( $13.0\% \pm 1.2\%$ ) showed in August 455 with the lowest water tableWTDgroundwater level of 3.1 m in 2019, whereas river water contributed the most <u>highest (47.1%  $\pm$  13.2%) to riparian groundwater in July with a higher water table groundwater level</u> of 1.8 m in 2021 (Figs. 3 and 9). The proportional contribution of the in-situ groundwater contributions to for riparian groundwater increased with the increasing distances from the riverbank during the observation periods, while the RWCs to riparian groundwater decreased significantly from D05 to D45 (p < 0.05) (Fig. 89). As shown in Table 460  $\frac{2}{2}$ , tThere was a significant increase of  $\frac{222}{Rn}$  activities activity in groundwater from D05 ( $\frac{494.5610.1}{2} \pm 107.5$ Bq/m<sup>3</sup>) to D45 (787.4  $\pm$  153.2 Bq/m<sup>3</sup>) ( $p \le 0.05$ ) (Table 1). The T<sub>res</sub> of recharged groundwater from river water that recharged by river to the underlying aquifer and/or riverbank increased from D05 ( $-0.09 \pm 0.090$  days) to D45 (0.15  $\pm$  0.13 days) (Table 21). These also indicated that the river recharged to-riparian deep strata was rapidly and frequently, particularly more evident significant in the plots closer to the riverbank.

#### 465 **3.<u>34</u>**Seasonal variations in RWC to riparian trees

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Significant differences in the seasonal RWCs to riparian *S. babylonica* were found between dry 2019 and wet 2021 (p < 0.05). As shown in Fig. 9, t<u>T</u>he proportional contributions of river water to <u>riparian *S. babylonica* trees</u> were significantly <u>more higher</u> in 2021 (mean of 23.8% ± 7.8%) than those in 2019 (mean of 16.8% ± 4.7%) (p < 0.05). In particular, riparian *S. babylonica* took up significantly more river water in July (35.2 ± 7.0%) and

- 470 November  $(29.0 \pm 5.0\%)$  in 2021 than that in July  $(15.4 \pm 1.7\%)$  and November  $(14.8 \pm 1.6\%)$  in 2019 (p < 0.001). The least absorption of river water for *S. babylonica* was  $17.7 \pm 2.7\%$  in 2021 (in September), which was larger than that of  $13.2 \pm 1.9\%$  in 2019 (in August). The most absorption of river water occurring in July in 2021 was also larger than that of  $24.2 \pm 3.0\%$  observed in June in 2019. Specifically, the most significantly monthly difference in the RWC to riparian *S. babylonica* trees between dry 2019 and wet 2021 was up to 19.8% (p < 0.001).
- 475 The monthly maximum RWC to *S. babylonica* trees was significantly higher in wet 2021 ( $35.2\% \pm 7.0\%$ ) compared to dry 2019 ( $24.2\% \pm 3.0\%$ ) (p < 0.05).

The riparian *S. babylonica* took up the most river water in July  $(35.2 \pm 7.0\%)$  in 2021, whereas the highest RWC to riparian trees occurred in June  $(24.2\% \pm 1.6\%)$  in 2019. The minimum river water uptake for riparian *S. babylonica* in 2021 was in September  $(17.7\% \pm 2.7\%)$ , while trees took up the least river water in August 2019

480 (13.2%  $\pm$  1.9%). Although the precipitation amount in rainy season was much higher than in drought season ( $p \le 0.001$ ), no significant difference in the RWC to riparian *S. babylonica* trees was observed between the rainy and drought seasons in a same year ( $p \ge 0.05$ ) (Figs. 2 and 9). The difference values of the RWC to riparian trees between the rainy (-4.0%)-and dry (-4.4%) seasons were not significantly different ( $p \ge 0.05$ ) in both 2019 (-4.0%) -and 2021 (-4.4%) (Fig. 9). These showed that there were no significantly seasonal variations in the

#### 485 <u>RWC to riparian trees in a same within a year (p > 0.05).</u>

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The water uptake of river water by riparian *S. babylonica* was significantly different among-between the three plots in 2019 and 2021 (p < 0.05), while no difference was observed between the three plots in 2021 (p > 0.05) (Fig. 10). Specifically, Tthe RWC to riparian trees decreased significantly by 6.9% from D05 (20.0%) to D45 (13.1%) in 2019 (p < 0.05), whereas there corresponding value reduced little by 2.6% (from 25.3% to 22.72%) along the distances was no significant difference in 2021 (p > 0.05) (Fig. 10).

#### 3.4-5 Relationships of between leaf $\delta^{13}C_3$ with RWCs to riparian trees and WTD

The seasonal leaf δ<sup>13</sup>C of riparian *S. babylonica* in three plots varied significantly in 2019 and 2021 (p < 0.05).</li>
The leaf δ<sup>13</sup>C of riparian *S. babylonica* trees in 2019 (-27.7 ± 1.0 ‰) was remarkably largersignificantly higher in 2019 (-27.7‰ ± 1.0 ‰) than that in 2021 (-29.7‰ ± 0.7 ‰) (p < 0.05) (Table 12). There was a significantly increase of the leaf δ<sup>13</sup>C from D05 (-28.8‰) to D45 (-27.0‰) in 2019 (p < 0.05), while no significant difference in the leaf δ<sup>13</sup>C was observed among between the different distances in 2021 (p > 0.05). The smallest lowest leaf δ<sup>13</sup>C value of riparian trees showed occurred on August 15 in 2019 and July 14 in 2021, before when intense

rainfall occurred in both years.

There was a significantly negative relationship between the RWCs to riparian trees and the WTD ( $R^2 = 0.57$ ;

500 p =< 0.00001) (Fig. 10a11a). The leaf δ<sup>13</sup>C of riparian S. babylonica was found to be negatively correlated with the RWCs to riparian trees (R<sup>2</sup> = 0.61; p =< 0.00001) but positively related to the WTD (R<sup>2</sup> = 0.37; p =< 0.0001) in linear functions (p < 0.001) (Fig. 10b-11b and c). These indicated that deeper WTD (2.7 ± 0.3 m) resulted in and lesslower RWCs to riparian S. babylonica and resulted in higher leaf-level WUE in the dry-drier year of 2019. In comparison, the riparian S. babylonica under relatively shallower WTD (1.7 ± 0.5 m) led to greater higher</li>
505 RWCs but smaller-lower leaf-level WUE in 2021. It seemed that the riparian S. babylonica could consume the least amount of river water for transpiration to achieve the maximum WUE when the WTD was 4.0 m.

#### **4** Discussion

### 4.1 <u>Advantages and limitations of MixSIAR model and the iteration method</u><del>River recharge to riparian deep soil water and groundwater</del>

510 The iteration method in combination with the MixSIAR model and stable water isotopes is particularly useful for separating and quantifying the proportional contributions of the indirect river water source to transpiration flux of riparian trees nearby a losing river. This integration of methods is superiormore accurate than to-previous studies (Alstad et al., 1999; Zhou et al., 2017; White and Smith, 2020), which only considered the-river water as a direct water source of riparian trees without considering their distances from the riverbank and extents of the 515 lateral roots. The primary advantage is that of the combined method is that it explicitly identifies the direct and indirect water sources of riparian trees according to the distance from the riverbank, the extents of lateral roots, and the process of river recharging riparian deep water. Both the trace plots and three diagnostic tests (i.e., Gelman-Rubin, Heidelberger-Welch, and Geweke) were used to Both the trace plots and three diagnostic tests are used to checkensure that the MixSAIR model has converged (Stock and Semmens, 2013). Moreover, the 520 MixSIAR model has explicitly considered the uncertainties in the isotopic values and the estimates of source contributions compared to the simpler linear mixing models (Stock and Semmens, 2013; Ma et al., 2016; Stock and Semmens, 2013). The strength of the newly proposed multi-iteration method is that it can determine the total contributions of the indirect river water source to riparian trees-nearby a losing river. The multi-iteration will not stops until there is no significant difference between the results of the last two iterations, which reduces the 525 <u>calculation errors of the RWC to riparian trees.</u> This study quantified the RWCs to riparian deep soil water within the capillary fringe and groundwater nearby a losing river. Approximately one third of the riparian groundwater was recharged by river water, while 46.5% of riparian groundwater was not exchanged with river water or other water sources (Fig. 8). A short residence time (no more than 0.28 days) of groundwater recharged by river water in 2021 indicated that there were rapid and frequent interactions between river water and riparian groundwater quickly but not mixing with water held tightly in the soil pores (Brooks et al., 2010; Evaristo et al., 2015; Allen et al., 2019). It was consistent with previous studies that the lateral seepage of river water or rising water table could briefly saturate riparian soils but not replace/flush immobile waters or isotopically homogenize different water pools (Sprenger et al., 2019).

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However, the riparian deep-water sources were identified using the water isotopic data collected between two campaigns (an interval of about one month). The riparian soil water movement was complex, icated and the water isotopes might not be uniform between two campaigns along the losing river. Assuming the isotopic uniformity over such a time interval may cause uncertainties in estimating the RWC to riparian deep water. In addition, we assumed that the contributions of old river water (before initial time (t-1)) to riparian in-situ deep water were 540 identical with those contributions of current river water (during the observation period between t-1 and t) to riparian in-situ deep water in this study. This could increasenduce some uncertainties on the estimations of the RWC to riparian deep water and the RWC to riparian trees. To minimize this issue, wWater samples need to be collected more frequently to quantify the contributions of river water to riparian deep water and trees. There was 545 a significant decline of water table (0.8 m) between June and August in 2019, while the water table increased by 1.3 m from June to November in 2021. Changes in water tables significantly affected the interactions between riparian deep water and river water. This study confirmed that the rising water tables stimulated riparian deep water to exchange with river water. The river water contributed significantly more to riparian deep water in wet 2021 with shallower WTD of 1.7 m than that in dry 2019 with deeper WTD of 2.7 m (Figs. 7 and 8). It agreed 550 well with the report of Wang et al. (2021) that the amount of surface water recharge to riparian groundwater increased with reducing WTD (ranging from 16.3 m to 2.6 m) in Baiyangdian wetland, China. The impacts of river water on riparian deep water weakened significantly with the increase of distance from the river. It could be inferred from the significantly declining RWCs to riparian deep water and increasing residence time of groundwater along the gradient of distance (p < 0.05) (Figs. 7 and 8; Table 2). A number of previous studies also indicated that the declining water table with distance resulted in weakening hydraulic connection between river water and riparian groundwater (Stellato et al., 2013).

#### 4.2 RWC to riparian trees and effects of the distance from the river on RWC

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## 560 **4.2 <u>RWC to riparian trees and effects of the distance from the river on RWC</u>The roles of river water in riparian tree water uptake**

In this study, deep-rooted riparian trees nearby the losing river were identified to use a small proportion of river water (less than 25%) for transpiration (Fig. 10). The small RWC to riparian trees may be caused by three non-565 exclusive processes: fFirstly, the lateral roots of riparian trees further than 5 m away from the riverbank rarely took up river water directly when their outer-projected edges of canopy (less than 5 m in our study) were out of reach of the river (Busch et al., 1992; Thorburn and Walker, 1994). Instead, they took up riparian deep soil water/groundwater recharged by river water, which was-likely to restricted the RWC to transpiration flux-of riparian trees. Secondly, the ecohydrologyical separation (Brooks et al., 2010; Evaristo et al., 2015; Allen et al., 570 2019; Sprenger et al., 2019) might result in large isotopic discrepancies between fast-moving water flow and immobile water for plant water uptake. Although the residence time of recharged groundwater from river water was extremely short (no more less than 0.28 days) (Table 1), only one third of riparian groundwater was replaced by the lateral seepage of river water (Fig. 9). This probably indicated that river water recharged mobile groundwater quickly but could not completely replace water held tightly in the soil pores (Brooks et al., 2010; 575 Evaristo et al., 2015; Allen et al., 2019). It was consistent with Sprenger et al. (2019) who found that the lateral seepage of river water or rising groundwater level could briefly saturate riparian soils but not entirely replace/flush immobile waters or isotopically homogenize different water pools. Thirdly, several recent studies showed that the tree species even phreatophytic/deep-rooted trees predominantly extended roots into fine pores to take up immobile soil water (Evaristo et al., 2015; Maxwell and Condon, 2016; Evaristo et al., 2019). As mentioned above,

- 580 the immobile water could not be completely replaced by infiltrating river water, which eventually resulted in a small contribution of river water to deep-rooted riparian trees. This ecohydrologicaly separation perspective that plant accessible water pools were separated from the fast-moving water can also be supported by the findings that no significant difference in RWCs to riparian trees between rainy and drought seasons was observed in both dry and wet years (Fig. 9). Because riparian *S. babylonica* trees preferred to rely on immobile water in fine soil pores
- 585 and they would not change the river water uptake patterns when the fast-moving precipitation input increased (Brooks et al., 2010; Sprenger et al., 2019).

In contrast, Alstad et al. (1999) found that riparian *Salix* trees nearby a losing river on the northeast side of the Rocky Mountain National Park, Colorado relied on rivers for approximately 80% of its transpiration-flux. It is probably due to the fact that only river water and precipitation were considered as potential water sources for

590 riparian Salix in their study. The RWC to riparian Salix trees calculated by Alstad et al. (1999) could be overestimated because it was supposed tolikely included all proportions of the indirect river water, in-situ soil water and in-situ groundwater contributions to riparian Salix trees. In fact, river water seeped into the saturated/vadose zone across the riparian riverbank and it was taken up by riparian trees indirectly in the form of river-recharged deep soil water/groundwater. In our study, we separated the contributions of indirect river water
 595 source (i.e., river-recharged deep soil water in the 80–170 cm layer and groundwater) for riparian trees. The

accurate quantification of the indirect RWC to deep-rooted riparian trees could help to determine the effect of riparian plant water use on river runoff along a losing river.

We observed substantially variations in the RWC to riparian trees at interannual (between two years) and spatial (between three distances from the riverbank) scales (Fig. 10). The RWC to riparian *S. babylonica* trees in wet 2021 was 1.4 times higher on average than in dry 2019 (Fig. 10). This is mainly because that the higher water tablegroundwater level in wet year induced higher RWC to riparian deep water, while riparian *S. babylonica* trees presented similar root architecture (i.e., phreatophyte) associated with similar water source proportions between dry and wet years (Figs. 10). Thus, the indirect RWC to riparian phreatophyte trees in wet year was higher than in dry year. Although there was no significant difference in the deep water (below the 80 cm layer) contributions to riparian trees between three plots, we observed substantial effect of the declining water tablegroundwater level with increasing distance from the riverbank on the decreased indirect RWC to riparian *S. babylonica* trees were generally

attributed to the various RWCs to riparian deep water rather than the water use patterns of riparian trees. Our result is in contrasts to a previous study by Qian et al. (2017) who reported a significant increase of the RWC to G. 610 biloba trees in response to the water tablegroundwater level decline. This discrepancy was ascribed to the fact that riparian G. biloba had a dimorphic root system and shifted their main water sources from shallow soil layer to deeper soil layer. Nevertheless, the potential root growth rate of riparian phreatophyte S. babylonica trees can reach 1-13 mm/day, which allows the riparian S. babylonica trees to remain in contact with a rising/declining water tablegroundwater level and keepmaintain constant water uptake proportions from deep strata below the 80 615 cm depth (Naumburg et al., 2005)."This study identified that deep rooted riparian trees nearby a losing river used a smaller proportion of river water (less than 25%), even though the water uptake from deep soils and groundwater recharged by river water were considered (Fig. 9). This finding was similar to the results reported on riparian trees nearby the perennial stream (Busch et al., 1992; Thorburn and Walker, 1994). It underlined that other water sources rather than river water played dominant roles in riparian tree water uptake. Global studies also concluded that 89% 620 of plant transpiration relied on precipitation during the growing season or that stored in the substrates from past season (Miguez Macho and Fan, 2021).

As the outer projected edge of canopy were out of reach of river, the lateral roots of riparian trees further than 5 m away from the riverbank could not directly absorb river water (Busch et al., 1992; Thorburn and Walker, 1994). The RWC to riparian trees mainly depended on the river water recharge to deep soil water and groundwater (less than 30%). The river water exchanged rapidly and frequently with riparian mobile deep soil water/groundwater, but rarely with bound water in fine pores (Brooks et al., 2010; Evaristo et al., 2015; Allen et al., 2019; Sprenger et al., 2019). Nevertheless, the riparian trees predominantly extended roots into fine pores to take up bound water (Evaristo et al., 2015; Maxwell and Condon, 2016; Evaristo et al., 2019). The mobile deep soil water/groundwater was alternative water source for riparian trees especially under drought conditions. These 630 discrepancies between fast moving river water seepage and immobile water for plant water uptake probably led to the small contribution of river water to riparian trees.

The temporal and spatial variations in the RWCs were significantly affected by various water tables in this study. Shallower water table could increase the river water recharge to riparian deep water which riparian trees mainly relied on. It was evident that riparian deep rooted *S. babylonica* absorbed significantly more river water in 2021 with 1 m shallower WTD than that in 2019 (Fig. 9). The decreasing RWCs along the gradient of distance

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in both dry and wet years also confirmed that the water table decline could make riparian phreatophyte trees less dependent on river water (Zhou et al., 2017). In contrast to the findings in this study, riparian dimorphic *G biloba* shifted from shallow soil layer to deeper soil layer with significantly increasing absorption of river water when the water table declined more than 1 m (Qian et al., 2017). This indicated that riparian dimorphic trees could increase the proportions of river water absorption in response to the water table decline, while the RWCs to riparian phreatophytic/deep rooted trees decreased with the water table decline.

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### 4.3 <u>Link between RWC/WUE/WTD and itsthe implications</u>Responses of WUE to riparian tree uptake of river water and WTD

The water uptake patterns of riparian S. babylonica trees generally remainfollowed the characteristics of a 645 phreatophyte. We observed that the leaf WUE of all S. babylonica trees across three plots in both dry and wet years were negatively correlated with the indirect RWC to riparian trees and positively related to the-WTD, respectively (Figs. 10, 11b, and 11c). These relationships are in-consistent with previous studies (Behzad et al., 2022; Cao et al., 2020; Ding et al., 2020; Behzad et al., 2022). Higher leaf WUE associated with lower RWC to riparian trees and lower groundwater levels are likely due to thatbecause the water stress restricts the stomatal 650 conductance and further reduces transpiration rate of riparian trees. Specifically, dry 2019 was characterized as higher water demand (indicated by higher VPD) and lower water availability compared to wet 2021, but the energy resource (indicated by net radiation) for riparian trees was similar between two years (Figs. S1 and S2). We supposargued that the water limitation rather than the energy limitation regulated the leaf-level stomatal conductance of riparian S. babylonica trees. The high water demands but low river water availability in dry year probably resulted in stomatal closure of riparian trees to minimize the water loss, which could eventually lead to 655 a decrease of transpiration rate and even photosynthetic rate (Fabiani et al., 2021; Behzad et al., 2022; Fabiani et al., 2021). Aguilos et al. (2018) further found that the water stress would enhance radiation-normalized WUE<sub>5</sub> because the lack of water availability induced a morestronger reduction in transpiration than photosynthesis. On account of With no difference in the average net radiation between dry and wet years, the lower river water 660 availability in the dry year probably resulted in an increase of the leaf WUE. It can be inferred that riparian S. babylonica trees took up more river water and probably showed a consumptive river-water-use pattern in the wet year compared to the dry year. This agreed well with previous studies that the woody plants showed lower leaf WUE and consumptive water-use patterns in rainy season, while they showed higher leaf WUE and conservative

water-use patterns with lower soil water availability in dry season (Horton and Clark, 2001; Behzad et al., 2022; 665 Cao et al., 2020; Behzad et al., 2022-Horton and Clark, 2001). However, consumptive river water taken up by riparian trees could result in a great loss of river waterthe consumptive river water use by riparian trees could result in an overconsumption of river water, which should be avoided in the riparian zone of a losing river restored by "ecological water"There was a balance and coordination between the soil water availability and river water absorption as well as leaf level WUE of riparian trees, because obvious differences in the RWCs and leaf  $\delta^{13}$ C 670 values were found in three plots between dry and wet years in this study (Fig. 9 and Table 2). The riparian S. babylonica grew more reliance on river water and the WUE significantly decreased with frequent precipitation and rising groundwater level in wet year (p < 0.05) (Figs. 9, 10a and 10b). This might be ascribed to that the highwater availability could stimulate riparian trees to maximize their transpiration rate and show a profligate water use strategy with growing water extraction from river. It agreed well with previous studies that woody plants 675 showed profligate water-use patterns and relatively lower WUE in rainy season, whereas they had higher WUE and conservative water use patterns as the soil water availability decreased in dry season (Horton and Clark, 2001; Cao et al., 2020). However, the profligate water use strategy of riparian trees could result in overconsumption of river flow, which indicated that rising water tables would not be recommended in this shallow WTD area. These leaf-level ecophysiological characteristics (i.e., water use efficiency and transpiration rate) of riparian trees 680 responded quickly to the changing water tables. It was evident that the WTD played a critical role in river water acquisition and WUE of riparian trees nearby losing rivers (Mensforth et al., 1994; Horton and Clark, 2001; Qian et al., 2017; Zhou et al., 2017). This study indicated that the increasing WTD linearly reduced the reliance of riparian trees on river water sources, mainly due to the diminishing water exchanges between river water and riparian groundwater with the water table decline 685 (Figs. 7, 8 and 10a). In comparison, the declining water table proportionally increased the leaf  $\delta^{13}C$  of riparian S. babylonica (Fig. 10b), which was consistent with previous studies that an exponential growth function existed between the leaf  $\delta^{13}$ C of riparian Salix gooddingii and WTD ( $\delta^{13}$ C = 2.76 – 24.78×exp<sup>=0.02WTD</sup>, 0 < WTD <10 m) (Horton and Clark, 2001). In this study, the 4 m of WTD seemed to be optimal to coordinate the riparian plantwater relations, when the riparian S. babylonica consumed the least amount of river water for transpiration and 690 showed highest WUE. This benefited for balancing the relationship between the riparian tree growth and river

flow reservation. Nevertheless, there were some controversial views that the WUE of plant species firstly

increased and then decreased with increasing WTD, and the maximum WUE occurred when WTD was 1.44 m (Xia et al., 2018) or 6 m (Antunes et al., 2018). The knee point of WTD was not observed in this study, suggesting that further investigations should be conducted under deeper water tables (> 4 m) to quantify the relationships among water tables, WUE and RWCs to riparian trees.

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The WTD played a critical role in the river water uptake of riparian trees near aby losing rivers (Mensforth et al., 1994; Horton and Clark, 2001; Qian et al., 2017; Zhou et al., 2017). We observed that the proportional contributions of the river water source to riparian trees decreased linearly in response to groundwater level decline, leading to a proportional increase in leaf WUE (Fig. 11a and b). It was consistent with Horton and Clark (2001) who found an exponential increase of the leaf WUE of riparian *Salix gooddingii* with increasing WTD. As mentioned above, we emphasized the key role of reduced water availability to account for the control of on decreasing transpiration rate onthus enhancing leaf WUE in this study. Nevertheless, there were some controversial views that the-leaf WUE of plant species increased firstly and then decreased with the-increasing WTD (Antunes et al., 2018; Xia et al., 2018). This could be due to the fact that riparian trees could tolerate reduced water availability only within a species-specific threshold, beyond which xylem cavitation and even crown mortality occurs (Naumburg et al., 2005). These indicated that the optimal WTD for plant species was related to the highest leaf WUE, under that condition plant species could consume lower-css transpiration-water for transpiration to maximize CO<sub>2</sub> assimilation (Antunes et al., 2018; Xia et al., 2018). The break point of WTD was not observed in this study (Fig. 11a and b). Further investigations need to be conducted under deeper water

710  $\frac{\text{table}\text{groundwater levels (WTD > 4 m) to optimize the WTD and riparian plant-water relations.}$ 

Our results have important implications for untangling the trade-offs between riparian tree water use and river runoff management. Considering various meteorological conditions, water resources and the leaf WUE of riparian trees, tThe proportion of the RWC to riparian trees has been compared between dry and wet years to investigate the effects of river water availability on the water use characteristics of riparian trees. The riparian *S*.

715 *babylonica* trees remainshowed the highest WUE and the lowest river water uptake proportion under the lowest groundwater level condition (with the WTD of 4 m). While tThe rising water tablegroundwater level would trigger riparian trees to show a consumptive river-water-use pattern, which eshould not be recommended in the revegetated riparian zone beside an ecological-water-recharged losing river. Therefore, the relationships between the RWC to riparian trees, leaf-level physiological characteristics (e.g., leaf WUE) and hydro-meteorological

720 <u>conditions are critical and helpful</u> to protect the revegetated riparian zones and maintain river runoff sustainability.

#### 4.4 Further scopes

Riparian trees could only indirectly absorb river water that merges into riparian deep water when they grow at a certain distance away from the riverbank. Inaccurate estimation of the RWCs to riparian trees nearby the losing rivers would be resulted once river water is identified as a separate source, but it can be resolved by the
newly developed iteration method in this study. When river water merging into riparian deep water is utilized by riparian trees, the iteration method could accurately separate and quantify the indirect contributions of river water to riparian trees. The iteration method has been proven to be available for quantifying the RWCs to those riparian trees growing under shallow WTD conditions along the losing river in this research. However, it still requires further improvements under more cases with deeper WTDs. Moreover, the proportions of old river water merging into riparian deep water before the initial time need more investigation by collecting prior water samples. It is evident that quantifying the relationships among the RWCs to riparian tree species, WTD and WUE provides critical insights into coordinating and balancing the river water conservation and riparian plant transpiration in losing rivers.

#### **5** Conclusions

In ∓this study, we presented a new iteration method together with the MixSIAR model and stable water isotopes (δ<sup>2</sup>H and δ<sup>18</sup>O) and the MixSIAR model-to separate and quantify the proportional indirect-contributions of the indirect-river water source-to riparian *S. babylonica* in dry 2019 and wet 2021 along a losing river in Beijing, China. It was found that ∓the infiltrating river water could quickly exchanged with riparian mobile water quickly(with a proportion of 24.1%) but not completely mixing with waters held tightly in the fine pores. Riparian trees nearby a perennial streamlosing river generally extended roots into fine pores to access the o-slow-movinginmobile water sources, The isotopic discrepancies between the fast-moving water flow and the immobile water taken up by the roots led to a small RWC (20.3%) to transpiration flux of riparian treesonly taking up a small amount of fast moving river water (20.3%). The water deficit in the dry year probably induced stomatal closure and larger reduction in transpiration compared to the photosynthesis of riparian trees, thus leading to an evident increase of the-WUE than in the wet year. More river water could be absorbed and lower leaf δ<sup>13</sup>C was resulted in wet year when the WTD was 1 m shallower than that in dry year. The RWCs to riparian trees decreased

along the distance away from the riverbank. Increasing WTD linearly reduced the reliance of riparian trees on river water sources (p < 0.001). It was found that The leaf δ<sup>13</sup>GWUE –showed a negatively correlation with the RWCs to riparian trees but was positively related to the WTD in linear functions (p < 0.001). Riparian S.</li>
babylonica trees maintained the highest WUE and the lowest river water uptake proportion under deep groundwater condition (with the WTD of 4 m) in this study. These suggested that rising water tablegroundwater level would promote triggered riparian trees to show a profligate water use strategy and increase the river water nequisitions uptake and show a consumptive river-water-use pattern, which wshould not be recommended for the water resources management of a losing river restored by ecological water. The maximum WTD of 4 m seemed to be optimal for riparian plant water relations, maintaining highest water use efficiency and minimizing the plant transpiration. This study provides valuable insights into riparian afforestation related to water use and healthy riparian ecosystem healthenhancement.

Data availability: The data that support the findings of this study are available from the corresponding authorupon request.

Author contributions: YL: Investigation, Methodology, Formal analysis, Writing - original draft, Writing - review & editing; YM: Methodology, Formal analysis, Conceptualization, Writing - review & editing; XFS: Supervision, Writing - review & editing, Project administration; QZ: Methodology. LXW: Writing - review & editing.

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

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Acknowledgements: This work was supported by the National Natural Science Foundation of China
(41730749) and the National Key R&D Program of China (2021YFC3201203). Sincere thanks go to Xue Zhang,
Yiran Li, Lihu Yang and Binghua Li for their assistance in experiments.

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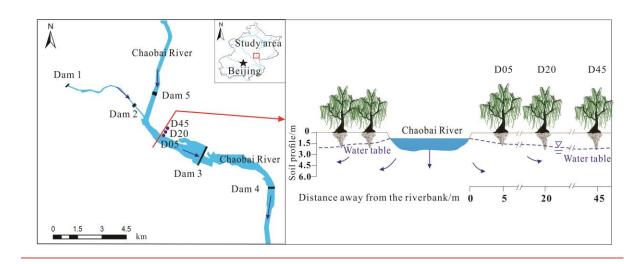
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**Figure 1:** Schematic diagram of the study area and the three sampling plots (D05, D20, and D45). D05, D20, and D45 are the plots at distance of 5 m, 20 m, and 45 m away from the

#### 925 riverbank, respectively.

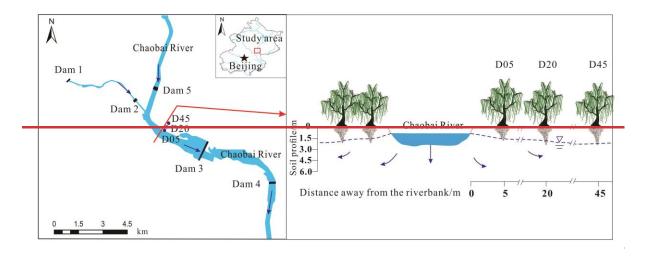
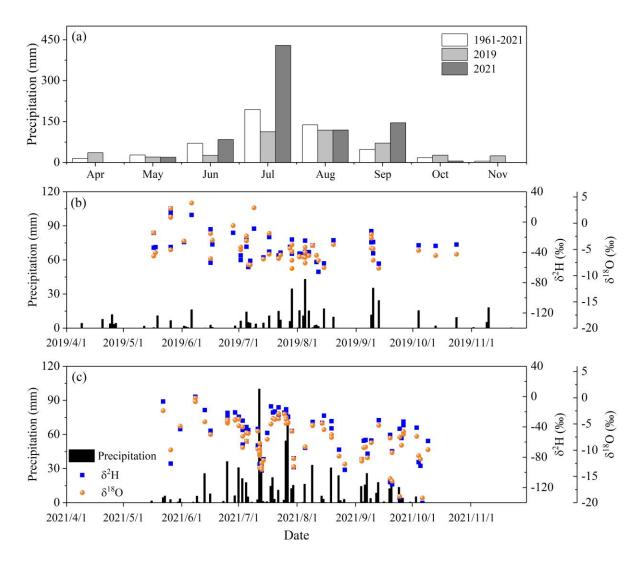


Figure 1: Schematic diagram of the study area and the three sampling plots (D05, D20, and D45). D05, D20, and D45 are the plots at distance of 5 m, 20 m, and 45 m away from the riverbank, respectively.



930 Figure 2: Changes in monthly average precipitation amount from 1961 to 2021 and monthly total precipitation amount for the observation years 2019 and 2021 (a), daily precipitation amount and precipitation isotopes during 2019 (b) and 2021 (c).

Figure 2: Changes in (a) average monthly precipitation between April and November from 1961 to 2021 and that in the observation years (2019 and 2021), and daily variations in the precipitation and their isotopic compositions in (b) 2019 and (c) 2021.

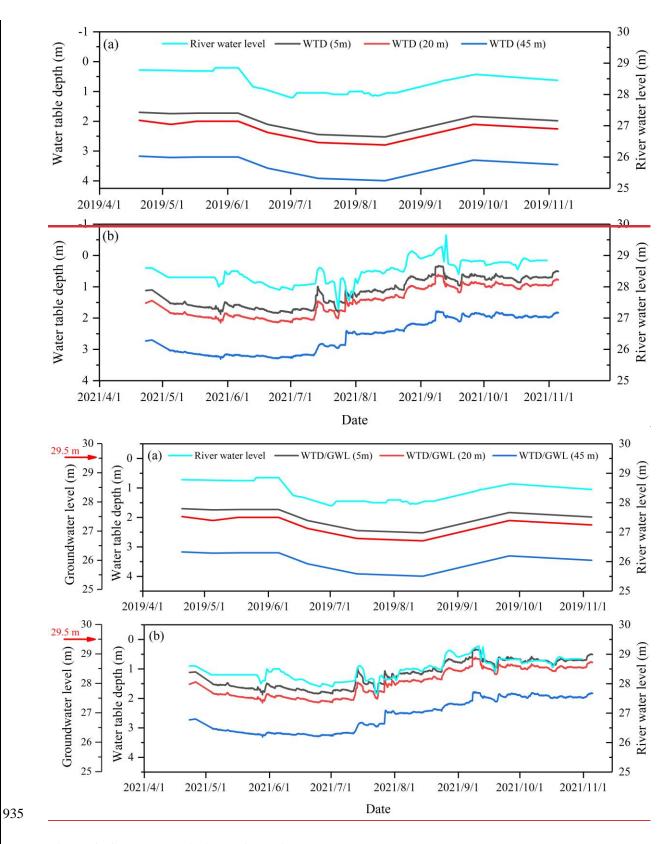


Figure 3: Seasonal variations of the river water level and <u>the depth to the water tablewater table</u> <u>depth depth (WTD)/ groundwater level (GWL)</u> at distances of 5 m, 20 m, and 45 m away from the riverbank during the observation period in <u>(a)-2019 (a)</u> and <u>(b)</u> 2021 (b). <u>The red arrow</u>

indicates the riparian ground surface level (29.5 m). The riverbed level is 26 m.

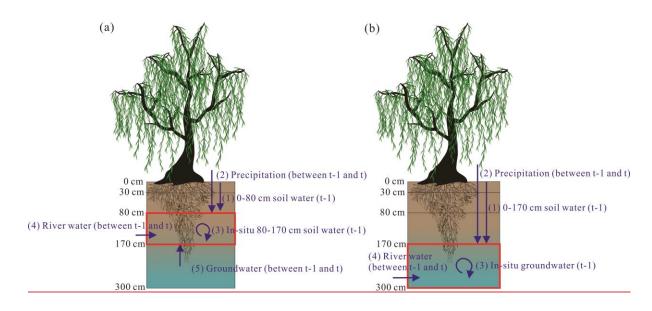


Figure 4: Schematic diagram for potential water sources of –riparian deep soil water in the 80–170 cm layer (a) and groundwater (b).

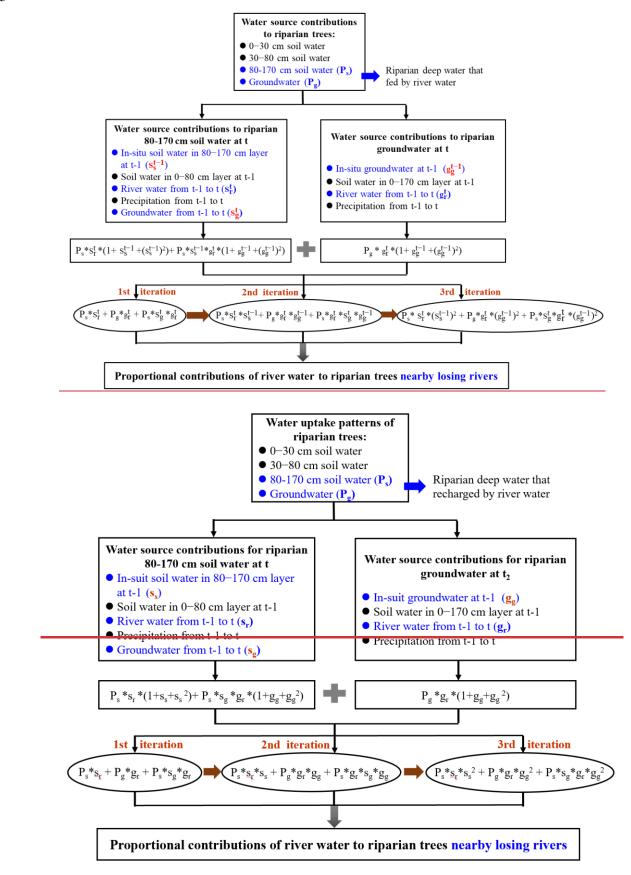


Figure 4: Flowchart for quantifying the proportional contributions of river water to riparian trees.

Figure 5: Flowchart for quantifying the proportional contributions of river water to riparian trees.

950 The P<sub>s</sub> and P<sub>g</sub> represent the contributions of riparian deep soil water in the 80–170 cm layer and groundwater to riparian trees, respectively. The s<sup>t-1</sup> and g<sup>t-1</sup> represent the proportional contributions of the old river water (before t-1) to riparian deep soil water in the 80–170 cm layer and groundwater, respectively. The s<sup>t-1</sup> s<sup>t</sup>, and s<sup>t</sup> represent the proportional contributions of in-situ soil water in the 80–170 cm layer at t-1, river water during t-1 to t, and groundwater during
955 t-1 to t for riparian deep soil water in the 80–170 cm layer at t, respectively. The g<sup>t-1</sup> and g<sup>t</sup> represent the proportional contributions of in-situ groundwater at t-1 and river water from t-1 to t for riparian groundwater at t, respectively.

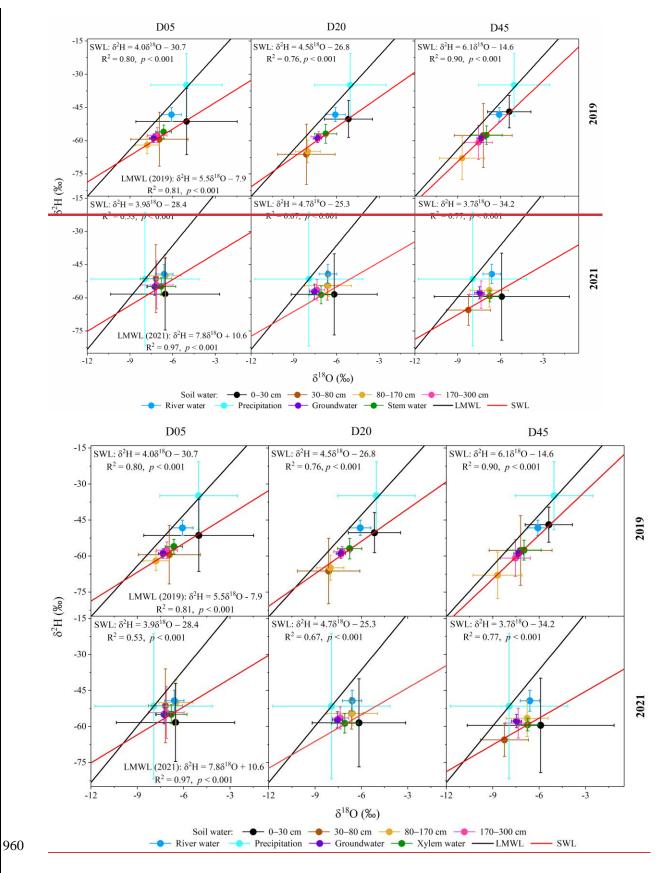


Figure <u>6</u>: Dual-isotope ( $\delta^2$ H and  $\delta^{18}$ O) biplots of different water bodies in the three plots (D05, D20, and D45) during the observation period infor the observation years 2019 and 2021. The

Local-local Meteoric-meteoric Water-water Line-line (LMWL) was fitted by the precipitation isotopic values of precipitation infor each year. The soil water line (SWL) was fitted by the soil water isotopes in all-the foursoil layers in-across the three plots (D05, D20, and D45) in-for each year. D05, D20, and D45 are the plots at distance of 5 m, 20 m, and 45 m away from the riverbank, respectively. The error bars indicate standard deviations.

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Figure 5: Dual-isotope ( $\delta^2$ H and  $\delta^{18}$ O) biplots of different water bodies in three plots (D05, D20, and D45) during the observation period in 2019 and 2021. The Local Meteoric Water Line (LMWL) was fitted by the isotopic values of precipitation in each year. The soil water line (SWL) was fitted by the water isotopes in all soil layers in the three plots (D05, D20, and D45) in each year. D05, D20, and D45 are the plots at distance of 5 m, 20 m, and 45 m away from the riverbank, respectively. The error bars indicate standard deviations.

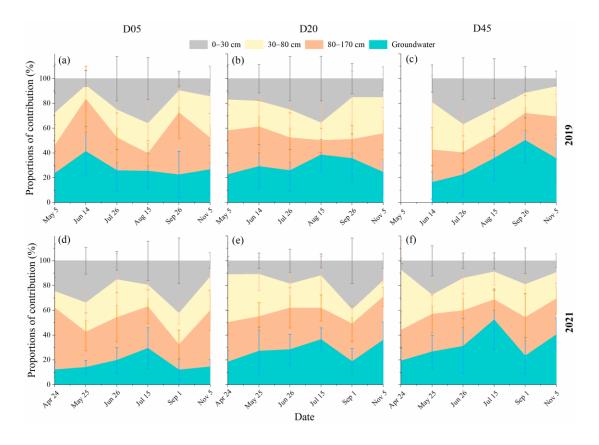


Figure 67: Seasonal water uptake patternvariations in the proportional contributions of the direct water sourcessoil water and groundwater to of riparian S. babylonicatrees in the three plots (D05, D20, and D45) during the observation period infor the observation years 2019 (a-c) and 2021 (d-f). D05, D20, and D45 are the plots at distance of 5 m, 20 m, and 45 m away from the riverbank, respectively. The error bars indicate standard deviations.

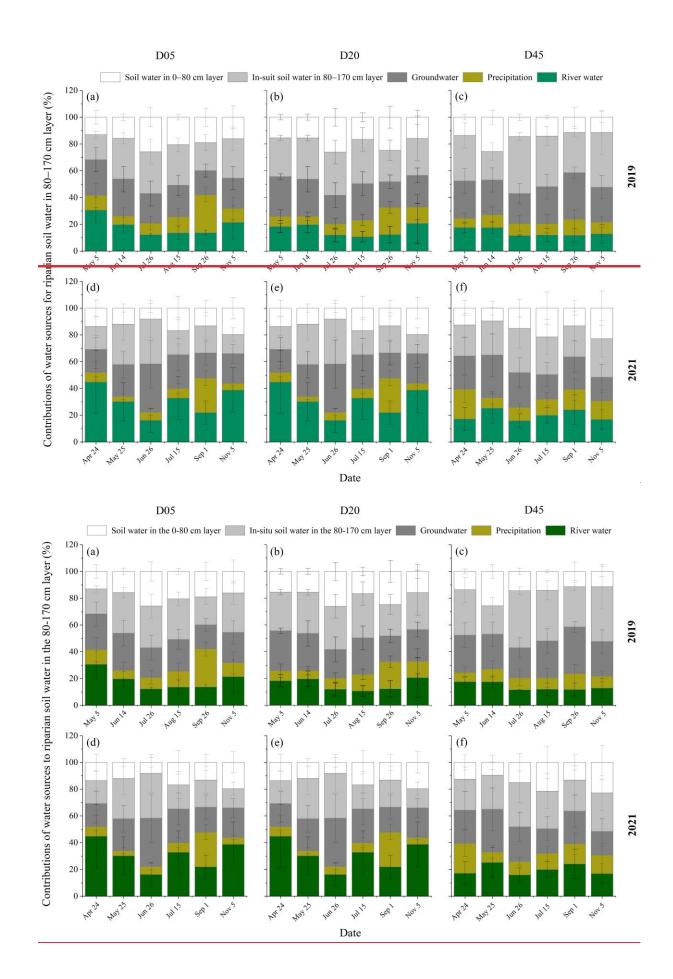
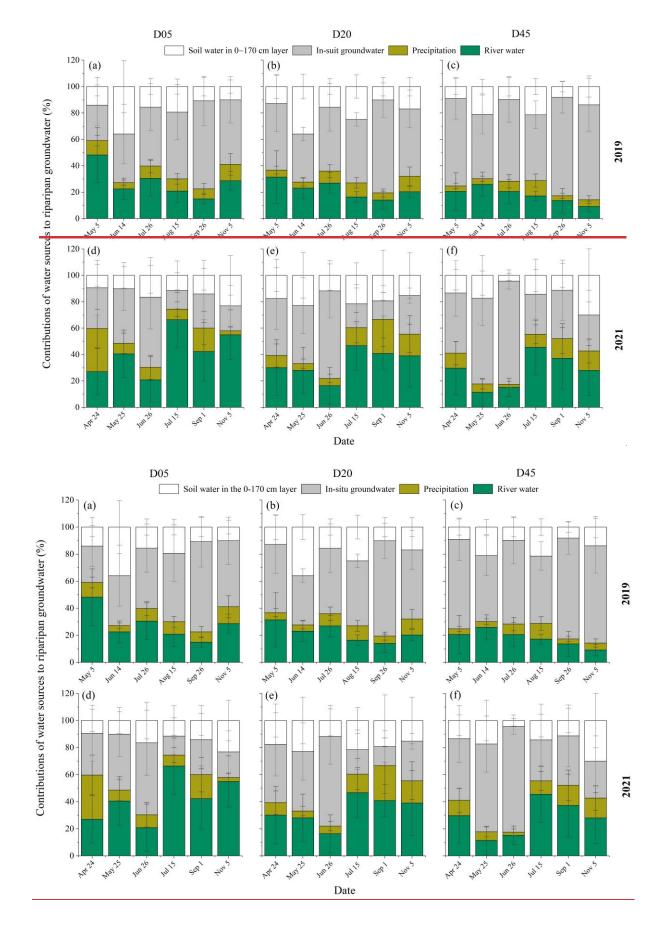


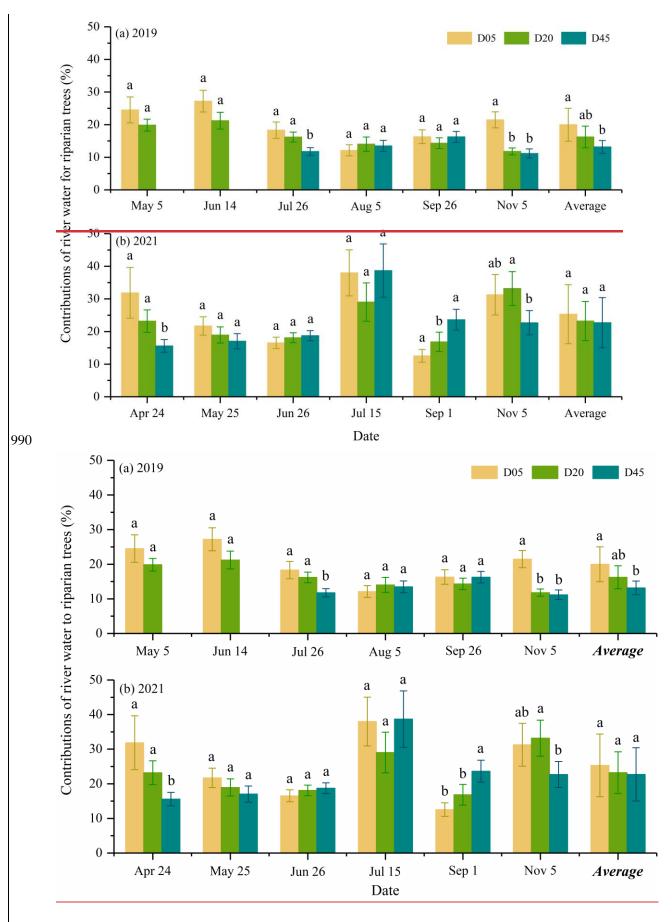
Figure 78: Seasonal variations in the different water source contributions to riparian deep soil water

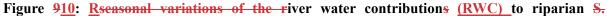
980 in <u>the 80–170 cm layer in the</u> three plots (D05, D20, and D45) <u>during the observation period in for the</u> <u>observation years</u> 2019 (a–c) and 2021 (d–f). D05, D20, and D45 are the plots at distance of 5 m, 20 m, and 45 m away from the riverbank, respectively. The error bars indicate standard deviations.





groundwater in <u>the</u> three plots (D05, D20, and D45)<u>during the observation period in for the</u> <u>observation years</u> 2019 (a–c) and 2021 (d–f). D05, D20, and D45 are the plots at distance of 5 m, 20 m, and 45 m away from the riverbank, respectively. The error bars indicate standard deviations.





babylonicatreesin the three plots (D05, D20, and D45) for each sampling campaign during theobservation period infor the observation years (a) 2019 (a) and (b) 2021 (b). The dD ifferent lettersrepresent show a significant differences in the river water contributions RWC to riparian S.babylonicatrees in between the three plots for each sampling campaign (p < 0.05). D05, D20, andD45 are the plots at distance of 5 m, 20 m, and 45 m away from the riverbank, respectively.

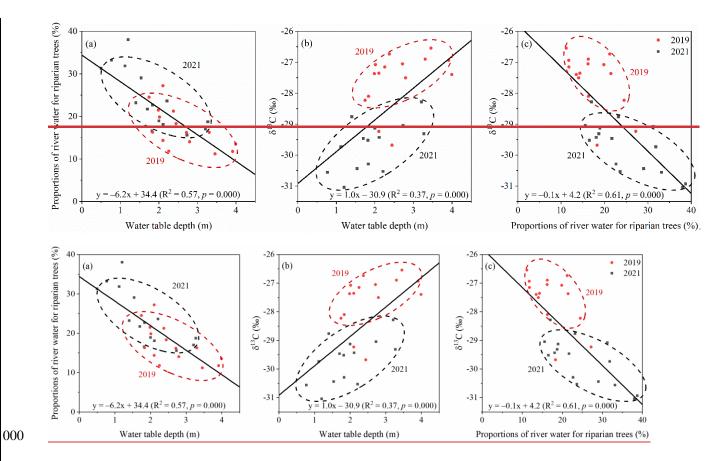


Figure 1011: Relationships between (a) the river water contributions proportions of river water contributions for to \_-riparian S. babylonicatrees and the water table depthwater table depth (a), between (b) the leaf  $\delta^{13}$ C values and the water table depthwater table depth (b), and between(c) the leaf  $\delta^{13}$ C values and proportions of river water contributions for river water contributions to riparian S. babylonicatrees (c).

|

Table 1: The <sup>222</sup>Rn values in river water, background groundwater and riparian groundwater in three plots (D05, D20, and D45), and the average residence time of recharged groundwater from river water (T<sub>res</sub>, day) in 2021. The background groundwater represents groundwater in aquifers more than 100 m away from the riverbank.

1010

	D:	Background	Riparian groundwater				
	<u>River water</u>	groundwater	<u>D05</u>	<u>D20</u>	<u>D45</u>		
$\frac{222}{\text{Rn value}}$	$\frac{610.1 \pm 212.3}{212.3}$	$\underline{7400\pm35.4}$	$610.1 \pm 107.5$	<u>763.3 ± 118.3</u>	$787.4 \pm 153.2$		
<u>T<sub>res</sub> (days)</u>	<u>0</u>	<u>Null</u>	<u>0</u>	$\underline{0.13\pm0.1}$	$\underline{0.15\pm0.13}$		

Notes: D05, D20, and D45 are the plots at distance of 5 m, 20 m, and 45 m away from the riverbank,

respectively.

	Leaf $\delta^{13}$ C value (‰)								
		2019							
	May 5	Jun 14	Jul 26	Aug 15	Sep 26	Nov 5	Mean	STD	
D05	-28.8	-29.2	-29.7	-30.4	-28.1	-27.4	-28.8	1.0	
D20	-27.1	-26.7	-27.1	-27.5	-27.4	-27.2	-27.1	0.2	
D45	Null	-27.2	-26.9	-27.4	-26.9	-26.5	-27.0	0.3	
	2021								
	Apr 24	May 25	Jun 26	Jul 14	Sep 1	Nov 5	Mean	STD	
D05	-29.7	-29.5	-29.5	-31.0	-29.5	-29.1	-29.7	0.6	
D20	-28.8	-29.1	-29.4	-30.4	-30.1	-30.3	-29.7	0.7	
D45	-29.0	-29.0	-29.4	-30.8	-30.1	-30.0	-29.7	0.9	

Table <u>2</u>: Leaf  $\delta^{13}$ C values of riparian *S. babylonica* in the three plots (D05, D20, and D45) during the observation period in 2019 and 2021.

Note: D05, D20, and D45 are the plots at distance of 5 m, 20 m, and 45 m away from the riverbank, respectively. STD represents standard deviations.-

 Table : The <sup>222</sup>Rn values in river water, background groundwater and riparian groundwater in three plots (D05, D20, and D45), and the average residence time of in 2021. The background groundwater represents groundwater in aquifers more than 100 m away from the riverbank.

	<b>D</b> ' (	Background-	Riparian groundwater				
	River water	groundwater	<del>D05</del>	<del>D20</del>	<del>D</del> 45		
<sup>222</sup> Rn value	$\frac{610.1 \pm 212.3}{10.1 \pm 212.3}$	7400	494.5 ± 107.5	$\frac{763.3 \pm 118.3}{2}$	<del>787.4 ± 153.2</del>		
( <del>Bq/m<sup>3</sup>)</del> T <sub>res</sub> (days)	θ	Null	$-0.09 \pm 0.09$	$0.13 \pm 0.1$	$0.15 \pm 0.13$		

Notes: D05, D20, and D45 are the plots at distance of 5 m, 20 m, and 45 m away from the riverbank, respectively.