



31 have extensively changed the hydrological characteristics of the riparian wetlands and might
32 accelerate carbon loss, which could further affect the GHG emissions.

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34 **Key words:** Riparian wetlands, Grasslands, Greenhouse gas, Spatial-temporal distribution, Impact
35 factor, Xilin River Basin

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39 1. Introduction

40 With the increasing impacts of global warming, the change in the concentrations of
41 greenhouse gases (GHGs) in the atmosphere is a source of concern in the scientific community
42 (Cao et al., 2005). According to the World Meteorological Organization (WMO, 2018), the
43 concentrations of carbon dioxide (CO₂), methane (CH₄), and nitrous oxide (N₂O) have increased
44 by 146%, 257%, and 122%, respectively, since 1750. Despite their lower atmospheric
45 concentrations, CH₄ and N₂O absorb infrared radiation approximately 28 and 265 times more
46 effectively at centennial timescales than CO₂ (IPCC, 2013). On a global scale, CO₂, CH₄, and N₂O
47 contribute 87% to the GHG effect (Ferrón et al., 2007).

48 Wetlands are unique ecosystems that serve as transition zones between terrestrial and aquatic
49 ecosystems. They play an important role in the global carbon cycle (Beger et al., 2010; Naiman
50 and Decamps, 1997). Wetlands are sensitive to hydrological changes, particularly in the context of
51 global climate change (Cheng and Huang, 2016). Moreover, wetland hydrology is affected by
52 local anthropogenic activities, such as the construction of reservoirs, resulting in gradual drying.
53 Although wetlands cover only 4–6% of the terrestrial land surface, they contain approximately
54 12–24% of global terrestrial soil organic carbon (SOC), thus acting as carbon sinks. Moreover,
55 they release CO₂, CH₄, and N₂O into the atmosphere and serve as carbon sources (Lv et al., 2013).
56 Wetlands are increasingly recognized as an essential part of the nature, given their simultaneous
57 functions as carbon sources and sinks. Excessive rainfall will cause an expansion in wetland areas
58 and a sharp increase in the soil moisture content (SMC), thus enhancing respiration,
59 methanogenesis, nitrification, and denitrification rates (Mitsch et al., 2009). On the contrary,
60 reduced precipitation or severe droughts will result in a decrease in water levels, causing the



61 wetlands to dry up. The accumulated carbon will be released back into the atmosphere through
62 oxidation. Due to the increasing impact of climate change and human activity, the drying of
63 wetlands has been widely observed in recent years (Liu et al., 2006); more than half of global
64 wetlands have disappeared since 1900 (Mitsch and Gosselink, 2007), and this tendency is
65 expected to continue in the future. The loss of wetlands may directly shift the soil environment
66 from anoxic to oxic conditions, and modify the CO₂ and CH₄ source and sink functions of wetland
67 ecological systems (Waddington and Roulet, 2000; Zona et al., 2013).

68 The Xilin River Basin in China is characterized by a marked spatial gradient in SMC. It is a
69 unique natural laboratory that may be used to explore the close relationships between the
70 spatiotemporal variations in hydrology and riparian biogeochemistry. Wetlands around the Xilin
71 River play an irreplaceable role with regard to local climate control, water conservation, the
72 carbon and nitrogen cycles, and husbandry (Gou et al., 2015; Kou, 2018). Moreover, the Xilin
73 River region is subjected to seasonal alterations in precipitation and temperature regimes, and
74 construction of the Xilin River Reservoir has resulted in highly negative consequences, such as the
75 drying of downstream wetlands, affecting riparian hydrology as well as microbial activity in
76 riparian soils. GHG emissions in riparian wetlands vary immensely. Understanding the
77 interactions between GHG emissions and hydrological changes in the Xilin River riparian
78 wetlands has thus become increasingly important. Moreover, it is necessary to estimate the
79 changes in GHG emissions as a result of wetland degradation at the local and global scales.

80 In this work, GHG emissions from riparian wetlands and adjacent hillslope grasslands of the
81 Xilin River Basin were investigated. GHG emissions, soil temperature (ST), and SMC were
82 measured in dry and wet seasons. The main objectives of this study were to (1) investigate the
83 temporal and spatial variations in CO₂, CH₄, and N₂O emissions from the wetlands in the riparian
84 zone, and examine the main factors affecting the GHG emissions, (2) compare the GHG emissions
85 from the riparian wetlands and different types of grasslands, and (3) evaluate the impact of
86 wetland degradation in the study area on GHG emissions.

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88 **2. Materials and methods**

89 **2.1 Study site**

90 The Xilin River is situated in the southeastern part of the Inner Mongolia Autonomous

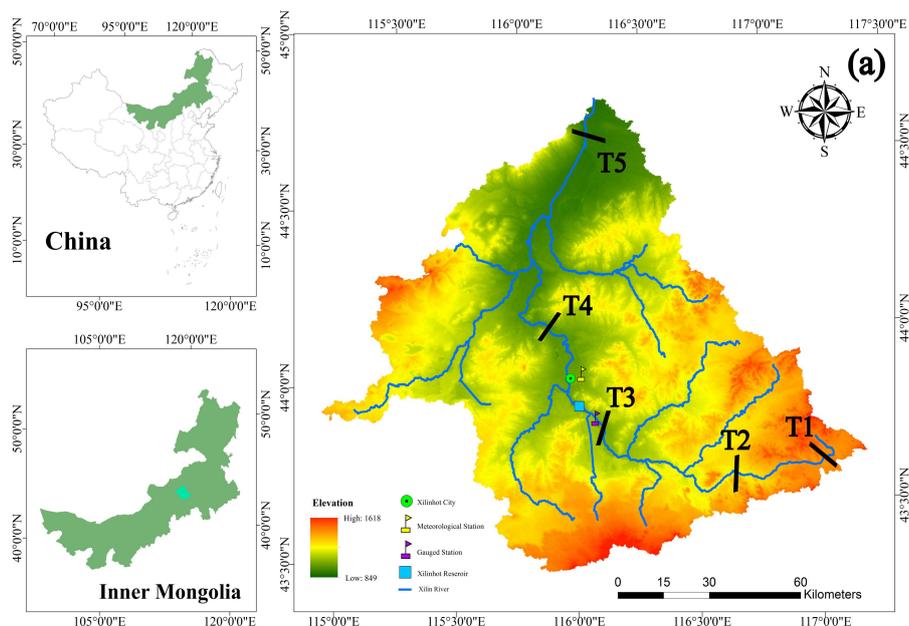


91 Region in China (E115°00'–117°30', N43°26'–44°39'). It is a typical inland river of the Inner
92 Mongolia grasslands. The river basin area is 10,542 km², the total length is 268.1 km, and the
93 average altitude is 988.5 m. According to the meteorological data provided by the Xilinhot
94 Meteorological Station (Xi et al., 2017; Tong et al., 2004), the long-term annual mean air
95 temperature is 1.7°C, and the maximum and minimum monthly means are 20.8°C in July and
96 –19.8°C in January, respectively. The average annual precipitation was 278.9 mm for the period of
97 1968–2015. Precipitation is distributed unevenly among the seasons, with 87.41% occurring
98 between May and September.

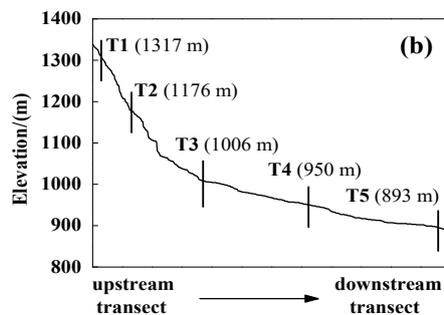
99 Soil types in the Xilin River Basin are predominantly chernozems (86.4%), showing a
100 significant zonal distribution as light chestnut soil, dark chestnut soil, and chernozems from the
101 northwest to southeast. Soil types in this basin also present a vertical distribution with elevation.
102 The chernozems are primarily soluble chernozems and carbonate chernozems, distributed at
103 altitudes above 1350 m with a relatively fertile and deep soil layer. Dark chestnut soil, boggy soil,
104 and dark meadow with high humus content are distributed between the altitudes of 1150 and 1350
105 m. Light chestnut soil, saline meadow soil, and meadow solonchak with low soil humus, a thin
106 soil layer, and coarse soil texture are distributed between the altitudes of 902 and 1150 m (Xi et al.,
107 2017).

108 **2.2 Field measurements and laboratory analyses**

109 In this study, five representative transects were selected as the primary measurement sites in
110 the entire Xilin River. Each transect cuts through the riparian wetlands near the river and hillslope
111 grasslands further away from it (Fig. 1).



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114 Fig. 1 (a) Location of the Xilin River Basin and distribution of five riparian-hillslope transects
115 (T1–T5). (b) Elevation details of each transect in the Xilin River Basin.

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117 The layout of the sampling points of each transect is shown in Fig. 2. Each sampling point for
118 T1–T5 was extended from the river to both sides, to the grassland on the slopes, using 5–7
119 sampling points for each transect and resulting in 24 points in total. The sampling sites on the left
120 and right banks were defined as L1–L3 and R1–R4 from the riparian wetlands to the hillslope
121 grasslands. As transect T3 was located on a much wider flood plain, none of its sampling points
122 were located on the hillslope grassland. The last transect (T5) was located downstream in the dry
123 lake and contained seven sampling points. They were defined as S1–S7, where S1, S2, and S7

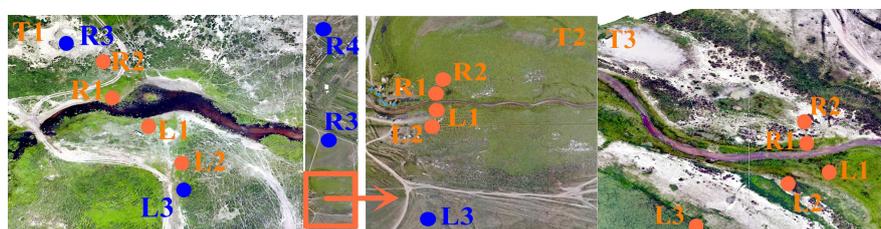


124 were located along the lake shore (the lakeside zone), and S3–S6 were located in the dry lake bed
125 (S3 and S4 in the mudbank, S5 in saline–alkali soil, and S6 in sand–gravel geology). Moreover,
126 characterizations for T1, T2, and T3 transects were the continuous river flow and T4 and T5
127 transects were the intermittent river flow.

128 The CO₂, CH₄, and N₂O emissions from each site were measured in August (wet season) and
129 October (dry season) in 2018 using a static dark chamber and the gas chromatography method.
130 The static chambers were made of a cube-shaped polyvinyl chloride (PVC) pipe (dimensions: 0.4
131 m × 0.2 m × 0.2 m). A battery-driven fan was installed horizontally inside the top wall of the
132 chamber to ensure proper air mixing during measurements. To minimize heating from solar
133 radiation, white adiabatic aluminum foil was used to cover the entire aboveground portion of the
134 chamber. During measurements, the chambers were driven into the soil to ensure airtightness and
135 connected with a differential gas analyzer (Li-7000 CO₂/H₂O analyzer, LI-COR, USA) to measure
136 the changes in the soil CO₂ concentration. The air in the chamber was sampled using a 60 mL
137 syringe at 0, 7, 14, 21, and 28 min. The gas samples were stored in the reservoir bag and taken to
138 the laboratory for CH₄ and N₂O measurements using gas chromatography (GC-2030, Japan). The
139 measurements were scheduled for 9:00–11:00 a.m. and 3:00–5:00 p.m.

140 ST was measured at depths of 0–10 cm and 10–20 cm with a geothermometer (DTM-461,
141 Hengshui, China). Plant samples were collected in a static chamber and over-dried in the
142 laboratory to obtain aboveground biomass (BIO). A 100 cm³ ring cutter was used to collect surface
143 soil samples at each site, which were placed in aluminum boxes and immediately brought back to
144 the laboratory to measure SMC and soil bulk density (ρ_b) using national standard methods
145 (NATESC, 2006). Topsoil samples were collected, sealed in plastic bags, and brought back to the
146 laboratory to measure soil pH, electrical conductivity (EC), total soil organic carbon (TOC), and
147 soil C:N ratio.

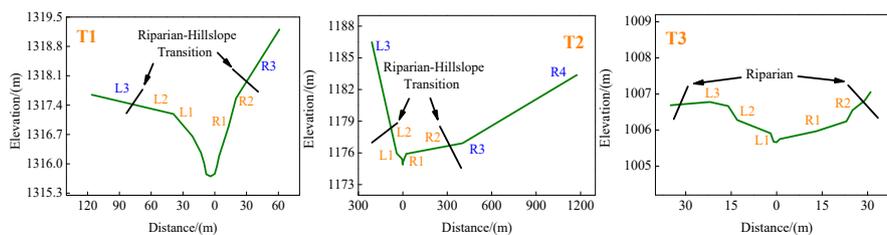
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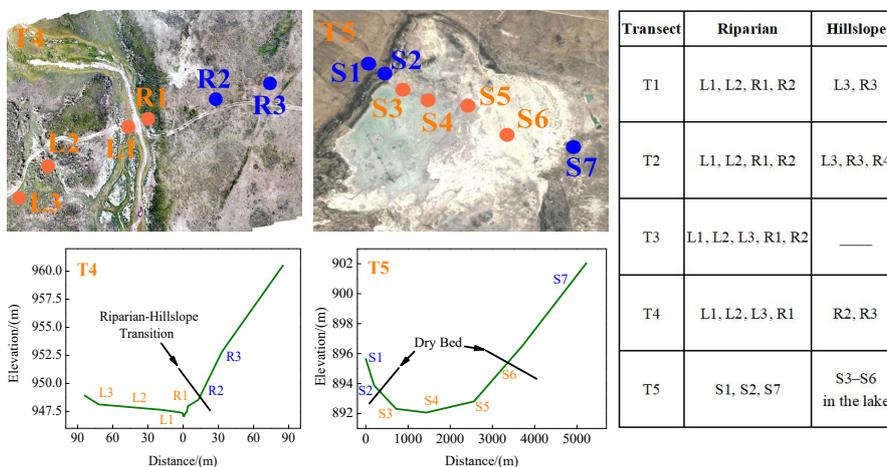
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Fig. 2 Distributions of sampling points in transects T1–T5 (The images are authors' own)

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Table 1. Physical and chemical properties of soils at various sites within each transect

Transect	Zone	Soil C:N	TOC (g·kg ⁻¹)	BIO (g)	ρ_b	pH	EC ($\mu\text{s}/\text{cm}$)
T1	Riparian	12.46 ± 0.91	30.16 ± 6.54	14.67 ± 5.44	1.28 ± 0.07	7.25 ± 0.62	154.71 ± 23.70
	Hillslope	11.41 ± 0.09	10.77 ± 4.72	6.70 ± 1.48	1.45 ± 0.03	7.22 ± 0.40	82.02 ± 16.37
T2	Riparian	11.70 ± 1.14	19.96 ± 5.71	24.76 ± 9.65	1.23 ± 0.05	8.95 ± 0.45	303.88 ± 102.16
	Hillslope	9.77 ± 0.88	14.87 ± 11.21	6.10 ± 3.19	1.38 ± 0.13	8.10 ± 0.55	162.97 ± 128.18
T3	Riparian	16.02 ± 3.74	25.16 ± 10.25	26.65 ± 40.64	1.09 ± 0.54	9.28 ± 0.72	1067.15 ± 813.13
T4	Riparian	12.52 ± 2.06	9.96 ± 1.25	11.97 ± 4.50	1.30 ± 0.08	8.84 ± 0.22	461.72 ± 314.27
	Hillslope	9.97 ± 0.50	9.65 ± 1.05	7.84 ± 2.48	1.30 ± 0.09	8.23 ± 0.14	118.5 ± 8.25
T5	Lake shore	63.74 ± 12.93	31.41 ± 6.55	5.48 ± 2.35	1.16 ± 0.10	9.88 ± 0.18	7320.87 ± 4300.03
	Dry lake bed	15.92 ± 4.71	6.35 ± 1.16	0	1.33 ± 0.09	9.41 ± 0.7	281.82 ± 162.73

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Note: Soil C:N - soil carbon-nitrogen ratio; TOC - total soil organic carbon; BIO - aboveground

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biomass; ρ_b - soil bulk density; pH - soil pH; EC - soil electrical conductivity



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159 **2.3 Calculation of GHG emissions**

160 The CO₂, CH₄, and N₂O emissions were calculated using Eq. 1 (Qin et al., 2016):

$$161 \quad F = \frac{V}{A} \times \frac{dc}{dt} \times \rho = H \times \frac{dc}{dt} \times \frac{M}{V} \times \left(\frac{273.15}{273.15 + t} \right), \quad (1)$$

162 where F denotes the CO₂, CH₄, and N₂O emissions (mg·m⁻²·h⁻¹), H is the height of the static
163 chamber (0.18 m), M is the relative molecular weight (44 for CO₂ and N₂O, and 16 for CH₄), V is
164 the volume of gas in the standard state (22.4 L·mol⁻¹), dc/dt is the rate of change of the gas
165 concentration (10⁻⁶h⁻¹), and T is the temperature in the black chamber (°C).

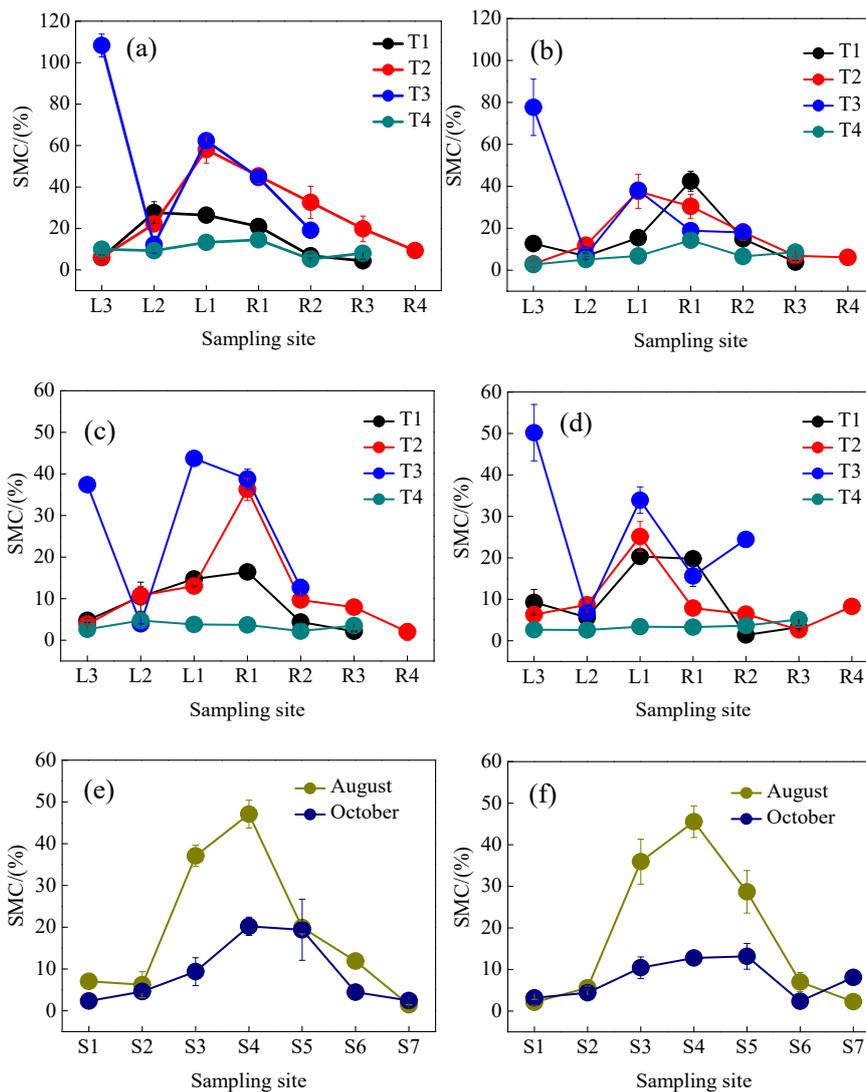
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167 **3. Results**

168 **3.1 Spatiotemporal patterns of SMC for each transect**

169 The spatiotemporal pattern of SMC is illustrated in Fig. 3. Similar variations were observed
170 between soil depths of 0–10 cm and 10–20 cm in the following order: wet season (August) > dry
171 season (October) and riparian wetlands > hillslope grasslands. The average SMCs in the
172 continuous river transects in the riparian zones (31.70% in August and 18.40% in October) were
173 higher than those in the hillslope grasslands (7.82% in August and 5.06% in October). During the
174 study period, the SMC changed as the distance from the river increased, and the highest value was
175 observed at the near-stream sites (L1 and R1). SMC fluctuations were low in the intermittent
176 transect compared to the upstream transects, with a mean value of 9.50% in August and 3.35% in
177 October in the riparian areas. The mean SMC in the hillslopes was 7.07% in August and 3.63% in
178 October. In transect T5, average SMCs at the center of the lake (29.15% in August and 11.52% in
179 October) were higher than those along the lake shore (4.12% in August and 4.18% in October).

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184 Fig. 3 Soil moisture contents (SMCs) at soil depths of 0–10 cm (SMC10) and 10–20 cm (SMC20)
 185 for transects T1–T5 in August and October: (a) SMC10 for T1–T4 in August. (b) SMC20 for
 186 T1–T4 in August. (c) SMC10 for T1–T4 in October. (d) SMC20 for T1–T4 in October. (e) SMC10
 187 for T5 in August and October. (f) SMC20 for T5 in August and October.

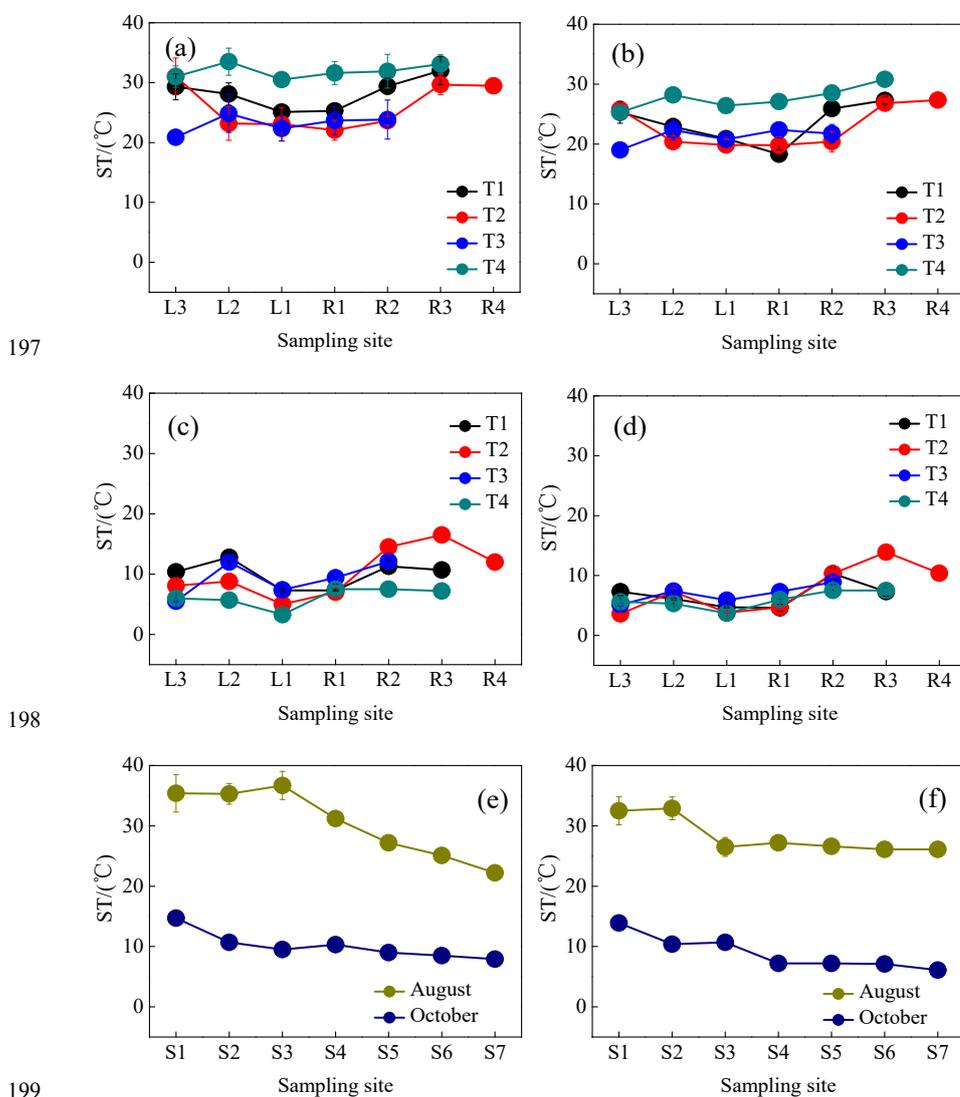
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189 3.2 Spatiotemporal patterns of ST in each transect

190 Spatiotemporal differences in ST during the entire observation period are displayed in Fig. 4.
 191 ST variations in August (mean value: 27.4°C) were noticeably higher than those in October (mean



192 value: 8.97°C). Moreover, ST for riparian sites (mean values: 26.0°C in August and 8.41°C in
 193 October) was slightly lower than that for the hillslope grasslands (mean values: 30.9°C in August
 194 and 10.3°C in October) for the 0–10 cm soil depth, with the exception of transect T5. Similar
 195 results were observed for the 10–20 cm soil depth.
 196



200 Fig. 4 Soil temperature (ST) at soil depths of 0–10 cm (ST10) and 10–20 cm (ST20) for transects
 201 T1–T5 in August and October: (a) ST10 for T1–T4 in August. (b) ST20 for T1–T4 in August. (c)
 202 ST10 for T1–T4 in October. (d) ST20 for T1–T4 in October. (e) ST10 for T5 in August and



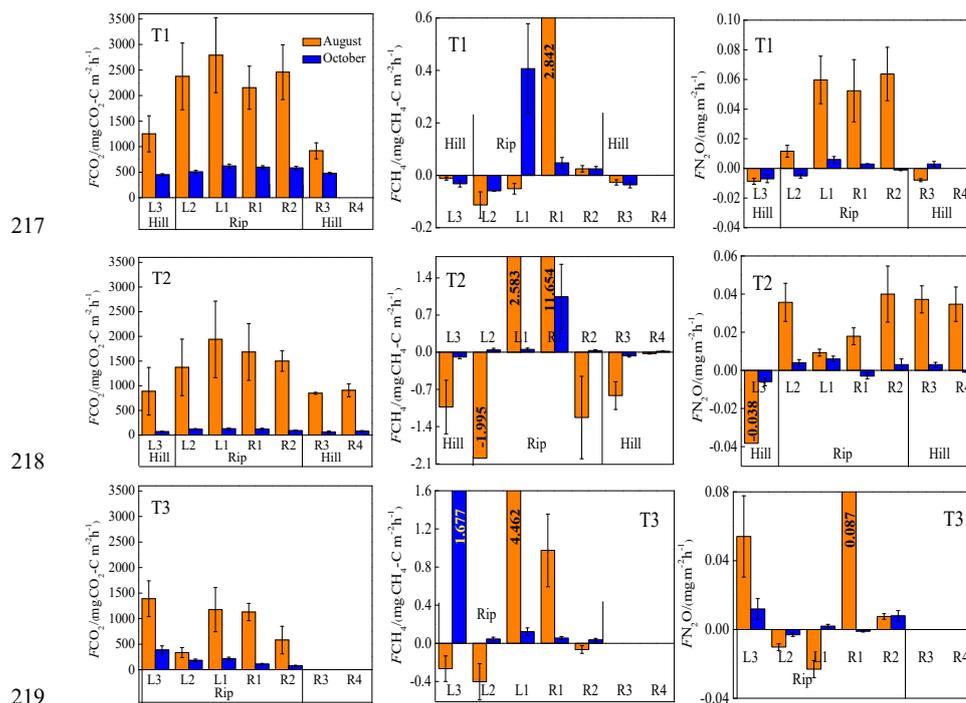
203 October. (f) ST20 for T5 in August and October.

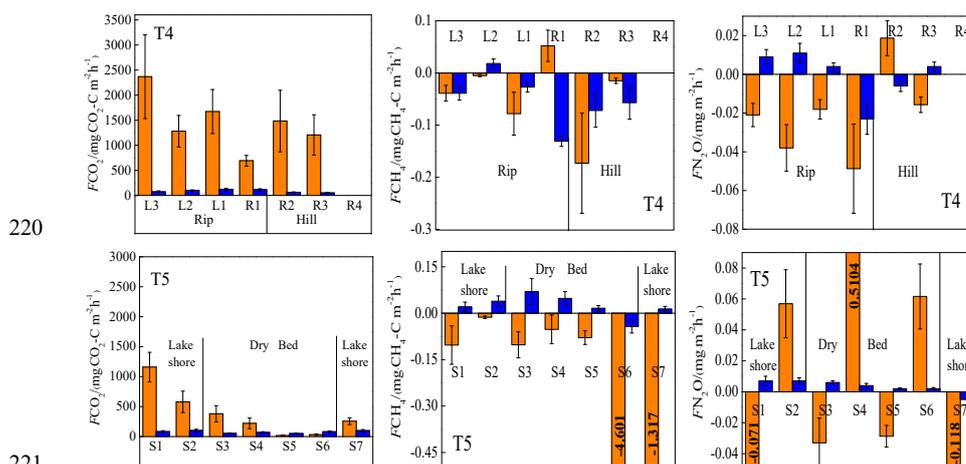
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205 **3.3 Spatiotemporal patterns of GHG emissions in each transect**

206 Figure 5 shows the spatiotemporal variations in GHG emissions in August (wet season) and
 207 October (dry season) in each transect. CO₂ emissions in each transect were higher in August than
 208 in October. The average emissions for the riparian wetlands of transects T1–T4 (1582.09 ± 679.34
 209 mg·m⁻²·h⁻¹ in August and 163.24 ± 84.98 mg·m⁻²·h⁻¹ in October) were higher than those for the
 210 hillslope grasslands (1071.54 ± 225.39 mg·m⁻²·h⁻¹ in August and 77.68 ± 25.32 mg·m⁻²·h⁻¹ in
 211 October). Higher CO₂ fluxes occurred in the riparian zones, while lower CO₂ fluxes were
 212 observed in the hillslope grasslands in continuous river transects (T1, T2, and T3). Transect T4
 213 exhibited lower CO₂ emissions in the riparian wetlands near the channel than at sites away from
 214 the channel. CO₂ emissions in transect T5 in August and October decreased from the lake shore to
 215 the lake center.

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Fig. 5 Spatiotemporal patterns of CO₂ (first column), CH₄ (second column), and N₂O

(third column) emissions (F) for each transect. Data are shown for August (orange) and October

(blue) and error bars are the standard deviations.

CH₄ emissions at the transects with continuous river flow (T1, T2, and T3) varied between August and October, except for T4 (characterized by intermittent river flow) and T5 (the dry lake). In August, the near-stream sites (L1 and R1) in T1, T2, and T3 were characterized as high CH₄ sources (average: $3.74 \pm 3.81 \text{ mg} \cdot \text{m}^{-2} \cdot \text{h}^{-1}$), but the sites located away from the river gradually turned into CH₄ sinks. Moreover, all the sites in transects T4 and T5 were sinks. CH₄ emissions (mean value: $0.2 \pm 0.45 \text{ mg} \cdot \text{m}^{-2} \cdot \text{h}^{-1}$) at the wetland sites were always lower in October than those in August. However, the sites on the hillslope grasslands served as CH₄ sinks (mean value: $-0.05 \pm 0.03 \text{ mg} \cdot \text{m}^{-2} \cdot \text{h}^{-1}$). In transect T5, CH₄ emissions revealed the opposite trend; a CH₄ sink was observed in August, but it was transformed into a CH₄ source in October.

Similar to the CO₂ and CH₄ emissions, N₂O emissions showed a distinct spatiotemporal pattern for all the transects. N₂O emissions in August were higher than those in October. These emissions were higher in riparian wetlands than in hillslope grasslands. Moreover, almost all the sites with continuous river flow were N₂O sources, while more than half of the sites with intermittent river flow were sinks.

3.4 Spatiotemporal patterns of GHG emissions in upstream and downstream



242 **areas**

243 Figure 6 shows the detailed spatial and seasonal distribution of GHG emissions in August and
 244 October in the longitudinal direction from the upstream (T1, T2, and T3) to the downstream areas
 245 (T4 and T5). The CO₂, CH₄, and N₂O emissions were calculated from the average values of the
 246 respective emissions in the wetlands and hillslope grasslands in each transect.

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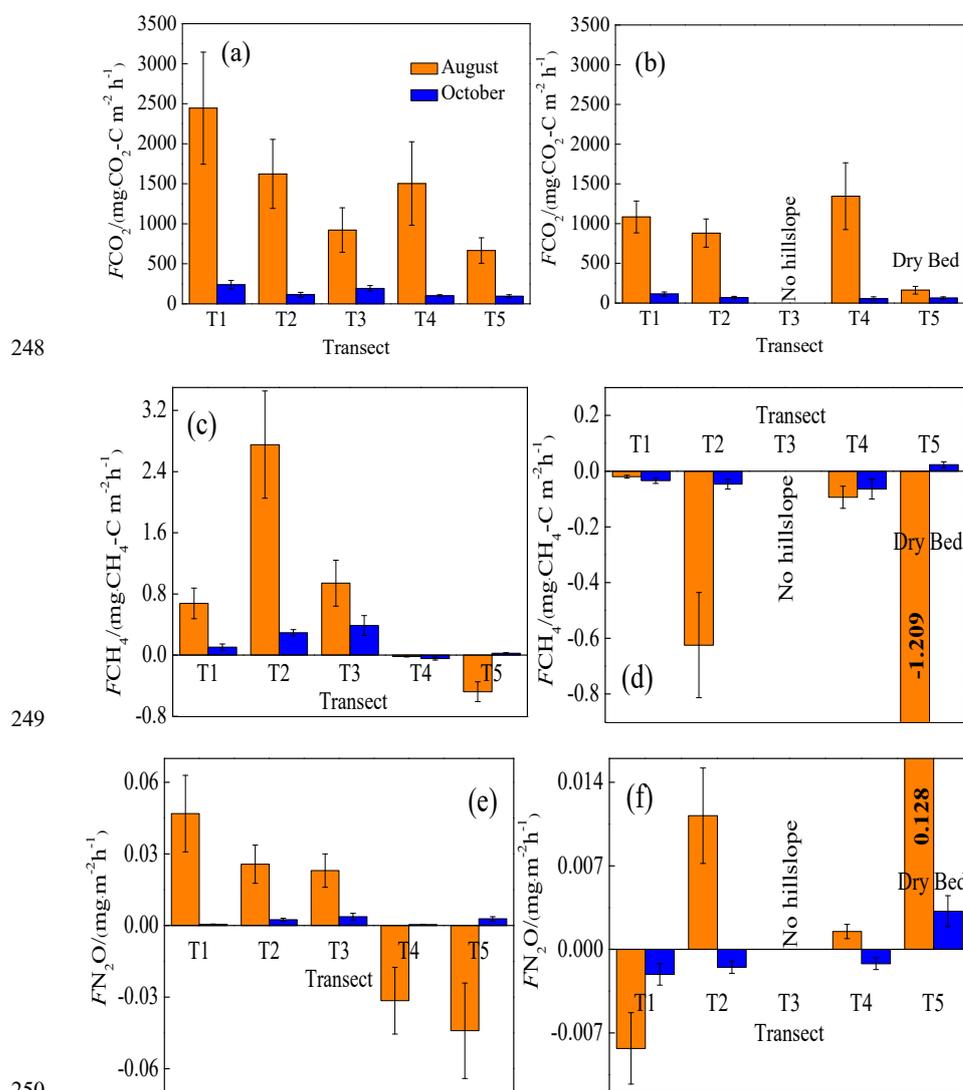


Fig. 6 Spatiotemporal patterns of CO₂ (first line), CH₄ (second line), and N₂O (third line) emissions (F) in the upstream (T1, T2, and T3) and downstream areas (T4 and T5). Data are



253 shown for the riparian wetlands (a, c, e) and hillslope grasslands (b, d, f) in August (orange) and
254 October (blue). Bars are the mean values for each transect and error bars are the standard errors.

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256 CO₂ emissions in riparian wetlands (Fig. 6(a)) in August decreased from 2444.69 ± 228.58
257 $\text{mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$ in the upstream area to $665.08 \pm 347.57 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$ downstream, and the
258 corresponding values for October were $238.12 \pm 48.20 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$ and $94.14 \pm 7.67 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$.
259 However, in hillslope grasslands (Fig. 6(b)), CO₂ emissions exhibited no significant seasonality
260 between upstream and downstream areas, with the mean values of $1103.40 \pm 190.44 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$
261 in August and $79.18 \pm 24.52 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$ in October. In addition, CO₂ emissions in transect T5
262 were lower for both months, with the averages of $162.83 \pm 149.15 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$ and 63.26 ± 12.40
263 $\text{mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$ in August and October, respectively. The upstream riparian zones exhibited higher
264 CO₂ emissions ($894.32 \pm 868.47 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$) than their downstream counterparts ($621.14 \pm$
265 $704.10 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$). However, mean CO₂ emissions showed no significant differences in
266 grasslands, averaging $524.16 \pm 450.10 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$ upstream and $508.06 \pm 534.77 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$
267 downstream.

268 CH₄ emissions showed a marked spatial pattern in the riparian zones from upstream to
269 downstream (Fig. 6(c)). The transects with continuous river flow were CH₄ sources in August and
270 October, with the average emissions of $1.42 \pm 3.41 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$ and $0.27 \pm 0.49 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$,
271 respectively, while those with intermittent river flow served as CH₄ sinks, with the corresponding
272 mean values of $-0.21 \pm 0.45 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$ and $-0.02 \pm 0.05 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$. Moreover, the hillslope
273 grassland sites in all transects were CH₄ sinks (Fig. 6(d)).

274 N₂O emissions in riparian wetlands (Fig. 7(e)) showed spatial patterns similar to those of
275 CH₄ emissions. In August, the transects with continuous river flow served as N₂O sources, with
276 the mean value of $0.031 \pm 0.031 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$, while those with intermittent river flow were N₂O
277 sinks with an average value of $-0.037 \pm 0.05 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$. In October, N₂O emissions occurred as
278 weak sources in the longitudinal transects, averaging $0.002 \pm 0.007 \text{ mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$. However, N₂O
279 emissions in hillslope grasslands did not show any spatial pattern (Fig. 7(f)).

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281 4. Discussion

282 4.1 Main factors influencing GHG emissions



283 4.1.1 Effects of SMC on GHG emissions

284 SMC constituted one of the main factors affecting GHG emissions in wetlands. In this study,
285 transects T1–T4 were characterized by a marked spatial SMC gradient (i.e., a gradual decrease
286 from the riparian wetlands to the hillslope grasslands, and from upstream to downstream (Fig. 3)).
287 The CO₂, CH₄, and N₂O emissions showed a similar trend. In Table 2, SMC10 is correlated with
288 CO₂ emissions ($P < 0.05$), SMC10 and SMC20 are significantly correlated with CH₄ emissions (P
289 < 0.01), and SMC10 and SMC20 are highly correlated with N₂O emissions ($P < 0.05$ and $P < 0.01$,
290 respectively). These results indicated the influence of wetland SMC on GHG emissions.

291 Typically, the optimal SMC values associated with CO₂ emissions in riparian wetlands range
292 from 40 to 60% (Sjögersten et al., 2006), and excessive SMC reduces soil gas transfer. On the
293 contrary, the SMC of hillslope grasslands is less than 10%, leading to a decrease in CO₂ emissions
294 compared to those in riparian zones (Moldrup et al., 2000). Similar results were obtained in our
295 study. The changes in CO₂ emissions in transect T5 were contrary to the change in the SMC likely
296 because other sediment properties for this transect were not conducive to the survival of
297 microorganisms (Table 1), and the increase in SMC did not increase the respiration activity of
298 microorganisms.

299 The largest CH₄ emissions were observed at the near-stream sites (i.e., L1 and R1) in T1, T2,
300 and T3, with the average SMC of 30.29%, while the SMC values at the other sites, which were
301 either weak sources or sinks, averaged at 14.57%. These results indicate that a higher SMC is
302 favorable for CH₄ emissions because a higher SMC denotes a soil in a reduced state, which is
303 beneficial for CH₄ production and inhibits CH₄ oxidation. A similar result was reported by Xu et al.
304 (2008). They conducted experiments of CH₄ emissions from a variety of paddy soils in China, and
305 showed that CH₄ production rates increased with the increase in SMC at the same incubation
306 temperature. Meng et al. (2001) also reported that water depth was the main factor affecting CH₄
307 emissions from wetlands. When the water level dropped below the soil surface, the decomposition
308 of organic matter accelerated, and CH₄ emissions decreased. If the oxide layer is large, the soil is
309 transformed into a CH₄ sink (Meng et al., 2011).

310 The N₂O fluxes showed a clear spatial pattern associated with the changes in SMC. When
311 SMC was below the saturated water content, N₂O emissions increased with the increase in SMC.
312 In contrast, when SMC was above the saturated water content, N₂O emissions gradually decreased



313 with the increase in SMC (Niu et al., 2017). In our study, N₂O emissions in the riparian wetlands
314 with a high SMC were higher than those in the hillslope grasslands, and N₂O emissions in the
315 upstream transects were higher than those downstream. These findings are consistent with those of
316 Liu et al. (2003), who showed that SMC is an essential factor affecting N₂O emissions.

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318 4.1.2 Effects of ST on GHG emissions

319 ST was another important factor affecting the CO₂ emissions in this study, as this parameter
320 was significantly correlated with CO₂ emissions ($P < 0.01$) (Table 2). The activity of soil
321 microorganisms increases with rising soil temperatures, leading to increased respiration, and
322 consequently higher CO₂ emissions (Heilman et al., 1999). Previous studies reported that ST
323 partially controls seasonal CO₂ emission patterns (Inubushi et al., 2003). Therefore, CO₂
324 emissions in August were significantly higher than those in October in this study.

325 CH₄ emissions showed a clear seasonal pattern because high summer temperatures improve
326 the activity of both CH₄-producing and -oxidizing bacteria (Ding et al., 2010). In this study, ST
327 was closely related to CH₄ emissions, especially in the near-stream sites, and CH₄ emissions in
328 August were 59 times higher than those in October. Conversely, seasonal fluctuations in CH₄
329 emissions at the downstream sampling sites were small. Table 2 indicates that the correlation
330 between CH₄ emissions and temperature is not significant because the production and emission of
331 CH₄ are significantly affected by water conditions. SMC showed a positive correlation with GHG
332 emissions. In addition, SMC affected ST to a certain extent, while the interactions between SMC
333 and ST had a mutual influence on CH₄ emissions. During the study period, the near-stream sites
334 (L1 and R1) maintained a super-wet state on the ground surface for a long time, which was
335 beneficial for the production of CH₄. However, the wetlands maintained a state without water
336 accumulation on the soil surface in the growing season, which was conducive to the oxidative
337 absorption of CH₄. SMC thus masked the effect of ST on CH₄ emissions.

338 Previous studies indicated that temperature is an important factor affecting N₂O emissions
339 (Sun et al., 2011) through primary mechanisms impacting the activities of the nitrifying and
340 denitrifying bacteria in soil. Table 2 shows that the correlations between N₂O emissions and ST10
341 and ST20 are poor ($P > 0.05$). This can be attributed to the wide suitable temperature range for
342 nitrification-denitrification and weak sensitivity to temperature. The average ST in August was



343 27.4°C, conducive to the growth of denitrifying microorganisms, while that in October was
344 8.97°C, and the microbial activity was generally low (Sun et al., 2011). Furthermore, ST
345 fluctuations were low both in August and October. Therefore, the effect of ST on N₂O emissions
346 was masked by other factors, such as moisture content.

347

348 4.1.3 Effects of BIO and soil organic matter on GHG emissions

349 CO₂ and CH₄ emissions were higher in the riparian wetlands than in the grasslands, mainly
350 because of greater vegetation cover. Typically, CO₂ emissions from riparian wetlands originate
351 from plants and microorganisms, with plant respiration accounting for a large proportion in the
352 growing season. Previous studies have shown that plant respiration accounts for 35–90% of the
353 total respiration in the wetland ecosystem (Johnson-Randall and Foote, 2005). Good soil
354 physicochemical properties and high soil total organic carbon (TOC) of riparian wetlands improve
355 the activity of soil microorganisms and plant root respiration. Table 2 shows that BIO is
356 significantly correlated with the CO₂ ($P < 0.05$) and CH₄ ($P < 0.01$) emissions. These results can
357 be attributed to the significant linear positive correlation between the respiration rate and plant
358 biomass (Lu et al., 2007). Higher plant biomass storage can achieve more carbon accumulation
359 during photosynthesis and higher exudate release by the roots. This, in turn, promotes the
360 accumulation of soil organic matter. Increased amount of organic matter stimulates the growth and
361 reproduction of soil microorganisms, ultimately promoting CO₂ and CH₄ emissions. Moreover,
362 plants act as a gas channel for CH₄ transmission, and a larger amount of biomass promotes CH₄
363 emissions, given the increased number of channels. In transect T3, high CO₂ emissions observed
364 at site L3 can be attributed to the relatively high levels of SMC, BIO, and soil nutrients, which
365 stimulate the microbial respiration rates.

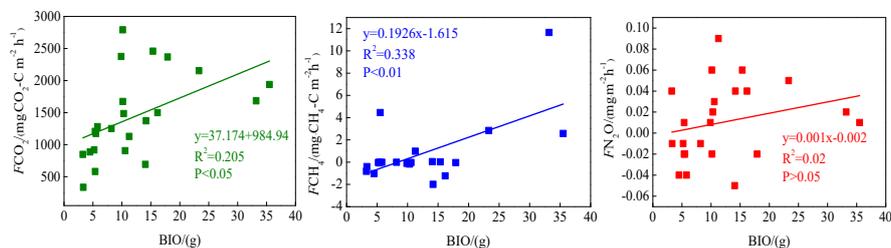
366 BIO had a weak correlation with N₂O emissions (Table 2), which indicates that plants
367 increase N₂O production and emissions, although this may not be the most critical factor. Previous
368 studies reported mechanisms wherein the plants can absorb N₂O produced in the soil through the
369 root system before releasing it into the atmosphere. Additionally, the root exudates of plants can
370 enhance the activity of nitrifying and denitrifying bacteria in the soil, ultimately promoting the
371 production of N₂O. Finally, oxygen stress caused by plant respiration can regulate the production
372 and consumption of N₂O in the soil, eventually affecting the conversion of nitrogen in the soil



373 (Koops et al., 1996; Azam et al., 2005).

374 Site L3 in transect T3 was covered by tall reeds, and its BIO was much higher than those of
375 the other sites; thus, the data for this site were excluded from the correlation analysis.

376



377

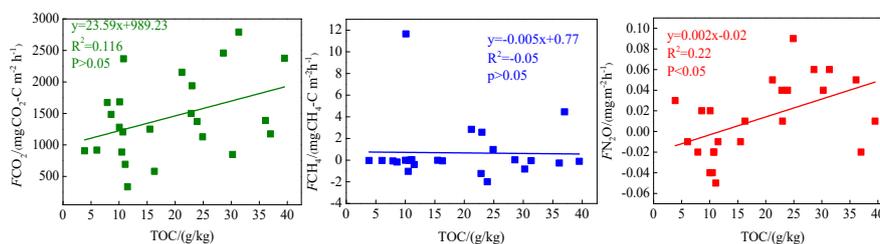
378 Fig. 7 Correlation between aboveground biomass (BIO) and GHG emissions (F)

379

380 The soil C:N ratio refers to the ratio of biodegradable carbonaceous organic matter and
381 nitrogenous matter in the soil, and it forms the soil matrix with TOC. TOC decomposition
382 provides energy for microbial activity, while the C:N ratio affects the decomposition of organic
383 matter by soil microorganisms (Gholz et al., 2010). The correlation results (Fig. 8) indicate that
384 TOC has a weak correlation with CO₂ emissions ($P > 0.05$), a poor correlation with CH₄ emissions
385 ($P > 0.05$), and a significant correlation with N₂O emissions ($P < 0.05$). Organic carbon provides a
386 carbon source for the growth of plants and microorganisms, which in turn boosts their respiration.
387 Moreover, as the abundance of heterotrophic nitrifying bacteria increases, soils become more
388 anaerobic, slowing down the growth of autotrophic nitrifying bacteria. This reduces the
389 nitrification rate, ultimately promoting N₂O release. Typically, low soil C:N ratios are favorable
390 for the decomposition of microorganisms, the most suitable range being between 10 and 12
391 (Pierzynski et al., 1994). Table 2 shows that N₂O emissions are significantly related to the soil C:N
392 ratios ($P < 0.05$), which means that denitrifying bacteria will use their own endogenous carbon
393 source for denitrification when the external carbon source is insufficient. Moreover, incomplete
394 denitrification leads to the accumulation of NO₂-N, which is conducive to the N₂O release. In this
395 study, all the sites in transects T1–T4 exhibited similar soil C:N ratios in the optimum range (Table
396 1), which is favorable for microbial decomposition. However, the soil C:N ratios in transect T5



397 were higher than those in the other transects, especially in the dry lake bed. Therefore, transect T5
 398 showed severe mineralization and a low microbial decomposition rate.
 399



400
 401 Fig. 8 Correlations between soil organic carbon (TOC) and GHG emissions (F)

402

403 Table 2. Correlations between CO_2 , CH_4 , and N_2O emissions and impact factors ($n = 62$)

GHG flux	ST10	ST20	SMC10	SMC20	TOC	ρ_b	C:N	pH	EC	BIO
CO_2	0.634**	0.592**	0.307*	0.216	0.393	-0.463**	-0.289*	-0.350**	-0.251*	0.491*
CH_4	-0.029	-0.051	0.346**	0.353**	-0.02	-0.129	-0.156	-0.127	-0.107	0.607**
N_2O	0.127	0.118	0.304*	0.356**	0.493*	-0.194	0.311*	0.137	0.504**	0.251

404 Note: 1. * and ** denote significant and highly significant correlations ($P < 0.01$ and $P < 0.05$),
 405 respectively.

406 2. ST - soil temperature, SMC - soil moisture content, ρ_b - soil bulk density, soil C:N - soil
 407 carbon-nitrogen ratio, pH - soil pH, EC - soil electrical conductivity, BIO - aboveground biomass

408

409 4.2 Riparian wetlands as hotspots of GHG emissions

410 The results of this study emphasized that CO_2 emissions in the riparian wetlands were higher
 411 than those in the hillslope grasslands owing to a variety of factors. ST is an important factor
 412 affecting GHG emissions. Mclain and Martens (2006) showed that seasonal fluctuations in ST and
 413 SMC in semi-arid regions have important effects on CO_2 , CH_4 , and N_2O emissions in riparian
 414 soils. Poblador et al. (2017) studied the GHG emissions in forest riparian zones and suggested that
 415 the difference in the CO_2 and N_2O emissions in these zones is affected by the spatial gradient of
 416 the regional SMC. In this study, the upstream riparian wetlands are characterized by higher TOC,
 417 lower soil C:N ratio, and abundant BIO than the hillslope grasslands (Table 1). These soil



418 conditions benefited the soil microbial activity, ultimately enhancing respiration as well as CO₂
419 emissions. However, CO₂ emissions in downstream areas were nearly identical to those in the
420 grasslands because the wetlands gradually evolved into grasslands after their degradation. The
421 N₂O emissions showed spatial patterns similar to those of the CO₂ emissions because the CO₂
422 concentrations were closely related to the processes of nitrification and denitrification, and high
423 CO₂ concentrations can promote the carbon and nitrogen cycles in soil (Azam et al., 2005; Baggs
424 et al., 2003), providing the substrate and energy required for the nitrification and denitrification
425 reactions. Moreover, soil respiration increases during soil denitrification (Liu et al., 2010;
426 Christensen et al., 1990). In this study, a weak correlation was observed between the CO₂ and CH₄
427 emissions in the riparian zones ($r = 0.228$), but CO₂ emissions were significantly correlated with
428 N₂O emissions ($r = 0.322$, $P < 0.05$). The soil became anaerobic in the riparian areas as the SMC
429 increased, and this was conducive to the survival of CH₄-producing bacteria and denitrification
430 reactions, eventually leading to an increase in CH₄ and N₂O emissions. Jacinthe et al. (2015)
431 reported that inundated grassland-dominated riparian wetlands were CH₄ sinks (-1.08 ± 0.22
432 $\text{kg}\cdot\text{CH}_4\text{-C ha}^{-1}\cdot\text{yr}^{-1}$), but Lu et al. (2015) indicated that grasslands were CH₄ sinks. In our study, a
433 marked water gradient across the transects led to the transformation of the soil from anaerobic to
434 aerobic soil, which changed the wetland function as a CH₄ source or sink. Therefore, during the
435 transition from the riparian wetlands to the hillslope grasslands, CH₄ emissions only appeared as
436 sources in the near-stream sites and sinks at other sites.

437 Further, we compared the GHG emissions of riparian wetlands and hillslope grasslands
438 around the Xilin River Basin with various types of grasslands (meadow grassland, typical
439 grassland, and desert grassland) in the Xinlingol League in Inner Mongolia (Table 3). The CO₂
440 emissions in August decreased in the following order: upstream riparian wetlands > downstream
441 riparian wetlands > hillslope grasslands > meadow grassland > typical grassland > desert
442 grassland. Moreover, the upper riparian wetlands acted as the source of CH₄ emissions, while the
443 downstream transects and grasslands served as CH₄ sinks. Similarly, except for the downstream
444 transects, N₂O emissions occurred as weak sources in different types of grasslands and upstream
445 riparian wetlands. The GHG emissions showed similar spatial patterns in October. Although these
446 estimates were made only in the growing season in August and the non-growing season in October,
447 our results suggest that riparian wetlands are the potential hotspots of GHG emissions. Thus, it is



448 important to study GHG emissions to obtain a comprehensive picture of the role of riparian
 449 wetlands in climate change.

450

451 Table 3. GHG emission fluxes of riparian wetlands and grasslands

Sample plot	GHG emissions in August ($\text{mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$)			GHG emissions in October ($\text{mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$)			Reference
	CO ₂	CH ₄	N ₂ O	CO ₂	CH ₄	N ₂ O	
Wetlands of upstream transects (T1, T2, and T3)	1606.28	1.417	0.031	182.35	0.272	0.002	This study
Wetlands of downstream transects (T4 and T5)	1144.15	-0.215	-0.037	98.13	-0.015	0.001	
Hillslope grasslands of all transects	1103.40	-0.246	0.001	79.18	-0.048	-0.002	Guo et al., 2017
Meadow grassland	166.39	-0.038	0.002	-	-	-	
Typical grassland	240.32	-0.042	0.037	-	-	-	
Desert grassland	107.59	-0.036	0.003	-	-	-	Zhang, 2019
Typical grassland	520.25	-0.102	0.007	88.34	-0.099	0.005	
Typical grassland	232.42	-0.090	0.004	-	-	-	Chao, 2019
Typical grassland	265.23	-0.185	0.005	189.41	-0.092	0.004	
Meadow grassland	553.85	-0.163	0.003	47.73	-0.019	0.011	Geng, 2004
Typical grassland	308.60	-0.105	0.002	70.25	-0.029	0.007	

452

453 4.3 Effects of riparian wetland degradation on GHG emissions

454 The hydrology and soil properties showed more evident differences among the transects
 455 because the downstream zone was dry all year due to the presence of the Xilinhot Dam (Fig. 1).
 456 The dam caused the degradation of the riparian wetlands, resulting in reduced GHG emissions.
 457 The average CO₂ emissions amounted to 1663 $\text{mg}\cdot\text{m}^{-2}\cdot\text{h}^{-1}$ in the riparian wetlands in the upstream
 458 transects (T1, T2, and T3), while the downstream transects (T4 and T5) recorded an average of



459 1084 mg·m⁻²·h⁻¹, 35% lower than the value in the upstream transects. The N₂O emissions from
460 the riparian wetlands were lower in the downstream transects.

461 The wetland degradation first resulted in the continuous reduction of SMC, affecting the
462 wetland carbon cycle processes. When the soil environment became less moist and more aerobic,
463 the CH₄ and N₂O emissions changed as the environment was transformed from a source to a sink,
464 and CO₂ emissions decreased. Table 1 shows that soil TOC in the upstream transects (average:
465 25.1 g·kg⁻¹) is higher than that in the downstream transects (average: 8.41 g·kg⁻¹). This result
466 indicates that wetland degradation caused the loss of the soil carbon pool and weakened the
467 wetland carbon source/sink function. These results are in agreement with those of Xia (2017).

468 The relatively low SMC and the aerobic environment were conducive to the mineralization
469 and decomposition of TOC. The degradation of plants in the wetlands led to the gradual reduction
470 of BIO. Ultimately, the plant carbon source input of the degraded wetlands decreased and the bare
471 land temperature increased due to the reduced plant shelter. This accelerated the decomposition of
472 TOC, leading to its decrease.

473 The degraded wetlands also caused soil desertification and salinization, leading to a decline
474 in the physical protection afforded by organic carbon and a reduction in soil aggregates. Thus, the
475 preservation provided by organic carbon declined. Soil TOC and SMC in the dry lake bed in
476 transect T5 were relatively high, but GHG emissions were very low along this transect because
477 soil pH values increased after the degradation of the lake soil, exceeding the optimum range
478 required for microorganism activity. The soil C:N ratio was very high, resulting in severe
479 mineralization and a low microbial decomposition rate, hence affecting the GHG emissions.

480

481 5. Conclusions

482 The riparian wetlands in the Xilin River Basin constitute a dynamic ecosystem. The present
483 spatial and temporal transfers in the studied biogeochemical processes were attributed to the
484 changes in SMC, ST, and soil substrate availability. Our simultaneous analysis of CO₂, CH₄, and
485 N₂O emissions from riparian wetlands and hillslope grasslands in the Xilin River Basin revealed
486 that the majority of the GHG emissions occurred in the form of CO₂. Moreover, our results clearly
487 illustrated a marked seasonality and spatial pattern of GHG emissions along the transects and in
488 the longitudinal direction (i.e., upstream and downstream). SMC and ST were two critical factors



489 controlling the GHG emissions. Moreover, abundant BIO promoted the CO₂, CH₄, and N₂O
490 emissions.

491 The riparian wetlands were the potential hotspots of GHG emissions in the Inner Mongolian
492 region. However, the degradation of wetlands transformed the area from a source to a sink for CH₄
493 and N₂O emissions, and reduced CO₂ emissions, which severely affected the wetland carbon cycle
494 processes. Overall, our study suggests that anthropogenic activities have significantly changed the
495 hydrological characteristics of the studied area, and will accelerate carbon loss from the riparian
496 wetlands and further influence the GHG emissions in the future.

497 **Author Contributions**

498 Xinyu Liu, Xixi Lu and Ruihong Yu designed the research framework and wrote the
499 manuscript. Xixi Lu and Ruihong Yu supervised the study. Xinyu Liu, Hao Xue, Zhen Qi,
500 Zhengxu Cao and Zhuangzhuang Zhang carried out the field experiments and laboratory
501 experiments analyses. Z.Z. drew GIS mapping in this paper. Tingxi Liu proofread the manuscript.

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507 **Competing interests**

508 The authors declare no conflicts of interest.

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