- 1 Methane emissions from a sediment-deposited island in a Lancang-Mekong
- 2 reservoir: Spatial heterogeneity and mechanisms
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## 12 Abstract

In dammed rivers, sediment accumulation creates potential methane emission hotspots, 13 which have been extensively studied in forebays. However, methane emissions from 14 sidebays remain poorly understood. We investigated methane emissions from a 15 sediment-deposited island situated in the sidebay of the Manwan Reservoir, Lancang-16 Mekong River. High methane emissions (maximum 10.4 mg  $h^{-1}m^{-2}$ ) were observed at 17 the island center, while a ring-like zone of low-to-negative methane emission was 18 discovered at the drawdown area around the island edge, whose flux varied between -19  $0.2-1.6 \text{ mg h}^{-1}\text{m}^{-2}$ . The ring-like zone accounted for 89.1 % of the island area, of which 20 9.1 % was a methane sink zone. Microbial processes in the drawdown area, regulated 21 by hydrological variations, were responsible for the low methane flux in this area. 22 23 Under reservoir operation, frequent water level fluctuations enhanced hyporheic exchange and created oxygen gradients along the hyporheic flow path. Dissolved 24 oxygen in hyporheic water decreased from 4.80 mg  $L^{-1}$  at the island bank edge to 0.43 25 mg  $L^{-1}$  at the center, which in turn decreased methanogen abundance for methane 26 production and increased methanotroph abundance for methane oxidation at the ring-27 like zone. This study quantified the methane emissions from sediment deposited islands 28 in the reservoir and helps to evaluate the global warming effects of hydropower systems. 29

### 30 **1 Introduction**

Natural rivers form continuous ecosystems, in which physical and chemical factors 31 32 drive biological processes from headwaters to river deltas (Butman and Raymond, 2011; Wilkinson et al., 2015). Along this continuum, rivers receive terrestrial organic carbon 33 (OC) and deliver it to the ocean at a global average rate of approximately 400–900 Tg 34 OC per year (Butturini et al., 2016; Seitzinger et al., 2005; Ran et al., 2013). In the past 35 two decades, many rivers have become intensively regulated by dams for a variety of 36 purposes, including improved navigation, water supply, flood control, and hydropower 37 38 production (Maavara et al., 2015). These engineering works decrease water velocity, converting rivers into a series of lentic reservoirs, where sediment accumulates in 39 forebays and sidebay islands (Maeck et al., 2013). Globally, the sediment accumulation 40 process has reduced the river-to-ocean flux of terrestrial OC by 26 % (Syvitski et al., 41 2005). 42

Methane is the second most important greenhouse gas, contributing approximately 43 44 18 % to total global warming effects (Smith et al., 2013; Wuebbles and Hayhoe, 2002). 45 Inland waters (lakes, rivers, and reservoirs) are significant sources of atmospheric methane, which is mainly released from anoxic sediment (Bastviken et al., 2011; Sobek 46 et al., 2012). After river damming, settling particles aggregate to form cohesive 47 sediment layers, which often become anoxic after oxygen is consumed but not 48 replenished through diffusive exchange (Rubol et al., 2013; Maeck et al., 2013). 49 Subsequently, large amounts of methane may be produced and released into the 50 atmosphere (Thornton et al., 1990; Maeck et al., 2013; Wilkinson et al., 2015), thereby 51

reducing the green credentials of hydropower. This issue has received considerable 52 attention in dammed rivers (Giles, 2006; Hu and Cheng, 2013). Maeck et al. (2013) 53 54 identified reservoirs as methane emission hotspots by comparing reservoir and riverine reaches, and estimated that global methane emissions have increased by 7 % due to 55 sedimentation in dammed rivers. In sidebays, the deposited sediments often created the 56 drawdown area under water level fluctuations, where water, heat, nutrients and 57 chemicals are exchanged and many biogeochemical reactions preferentially occur 58 (Tonina and Buffington, 2011; Cardenas and Markowski, 2010), potentially emitting 59 60 large amounts of greenhouse gases. Previous studies have mainly focused on methane emissions from dam forebays (Yang et al., 2013; DelSontro et al., 2010; DelSontro et 61 al., 2011), while the understandings of methane emissions from sediments deposited in 62 63 sidebays remain poor.

In reservoirs, the water level decreases during hydropower production, and increases 64 with the inputs from inflows during the stagnant hydropower production. As a result, 65 66 the water level frequently fluctuates following hydropower production demands, which enhances hyporheic exchange by driving water flow in and out of the drawdown area 67 (Tonina and Buffington, 2011; Hucks Sawyer et al., 2009). This may lead to changes of 68 oxygen conditions in the interior of the drawdown area. Zarnetske found a redox 69 gradient along the hyporheic flow paths in a third-order stream in the Willamette River 70 basin, USA (Zarnetske et al., 2011a). Methane from sediments is mainly produced by 71 72 methanogens, and is consumed by methanotrophs (Borrel et al., 2011). Methanogens and methanotrophs belong to anaerobic and aerobic microbes, respectively (Nazaries et 73

al., 2013). The oxygen conditions in sediments have been found to manipulate these
microbial processes and methane emission (Chamberlain et al., 2016; Shi et al., 2018).
Hence, we suppose the changes of oxygen conditions in sediments may alter methane
emission scheme on sediment-deposited islands in reservoir sidebays.

In this study, methane emissions from a sediment-deposited island were investigated in the sidebay of Manwan Reservoir, Lancang-Mekong River. Monitoring wells were established to probe hyporheic exchange and oxygen gradients across the island. Methanogen and methanotroph abundances in the sediment were analyzed using quantitative polymerase chain reaction (qPCR) to reveal the associated molecular mechanism. The objective of this study was to explore methane emission patterns from sediment-deposited zones in the reservoir sidebay and the underlying mechanisms.

85

#### 86 2 Methods

#### 87 **2.1 Study area**

88 The Lancang-Mekong River is a trans-boundary river in Southeast Asia and the tenth-89 largest river in the world, which originates from the Tibetan Plateau in China, continues into Myanmar, Lao PDR, Thailand, Cambodia and Viet Nam, and discharges into the 90 South China Sea. It has a length of approximately 4909 km, a total watershed area of 91 795,000 km<sup>2</sup> (Mekong River Commission, 2018). Due to rich hydropower resources, 92 cascade dams have been built on the mainstream of the Lancang-Mekong River. After 93 impoundment, about 33 sediment-deposited islands of  $4.3 \times 10^5$  m<sup>2</sup> formed in the 94 reservoir sidebay in the upstream section of the Lancang-Mekong River in China, of 95

which 81.8% are located at the convex bank, and 18.2% at the concave bank (Fig. S1). 96 This study selected the widely distributed type of island at the convex bank for 97 investigation, which is located in the Manwan Reservoir (24°43'44" N, 100°23'5" E). 98 The studied island has an oval shape with a surface area of  $1.3 \times 10^4$  m<sup>2</sup> (Fig. 1). 99 Manwan dam is the first-built one (completed in 1993) in the upstream section of the 100 Lancang-Mekong River in China, with a height of 132 m. The Manwan Reservoir has 101 a normal water level of 994 m, a total storage capacity of  $5.0 \times 10^8$  m<sup>3</sup>, an installed 102 capacity of  $1.5 \times 10^6$  kW and hydrological residence time of 0.78 a. Manwan has a 103 104 subtropical plateau monsoon climate, and the temperature features no distinct seasons. Under water level fluctuation induced by reservoir operation, the island bank is 105 frequently flooded (Fig. S2). 106

## 107 **2.2 Monitoring wells**

Ten monitoring wells were installed in the island bank at 0.5 (W1), 1.5 (W2), 3.5 (W3), 108 6.5 (W4), 10.5 (W5), 15.5 (W6) 20.5 (W7), 25.5 (W8), 30.5 (W9), and 35.5 m (W10) 109 from the waterline, respectively (Fig. S2). The wells were 90-mm diameter perforated 110 polyvinylchloride pipes, reaching a depth of 2.0 m below the ground surface. To prevent 111 flooding, the wells were extended 2.0 m aboveground. Due to hydropower production, 112 the reservoir runs in a pseudo-periodic hydrological regime with cyclic water level 113 fluctuations. Here, we monitored a complete cycle of water level fluctuation within 115 114 h. Water levels were measured every 10 min from 11 to 16 September 2016 using 115 automated water level recorders (U2000101, OneSetHoBo, USA), which were mounted 116 at the bottom of W5, W7, W10, and the reservoir (Fig. S3). 117

Instantaneous lateral fluid fluxes (q) across the island bank per unit length were
calculated following the Darcy Eq. (1) (Gerecht et al., 2011; Hucks Sawyer et al., 2009):

$$q(t) = -Kb \cdot \left[\frac{\vartheta h(x,t)}{\vartheta x}\right] \tag{1}$$

where *Kb* is sediment transmissivity, m d<sup>-1</sup>; *h* is hydraulic head, m; *x* is distance, m; and *t* is time, d. A positive *q* value indicates flow from the reservoir to the island. The island *Kb* was 0.99 m d<sup>-1</sup>, which was measured according to Philip (1993).

## 124 **2.3 Sampling and physicochemical analysis**

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After water level receded at the monitoring time of 100 h, groundwater (100 ml) was 125 126 carefully sampled in triplicate from each monitoring well with a portable peristaltic pump (SC-1/253Yx, Chongqing Jieheng Peristaltic Pump Co., Ltd., China), and then 127 filtered *in situ* using portable syringe filters for water DOC analysis. Sediment (5 g) 128 129 was synchronously collected in triplicate from 10 cm below the surface adjacent to each well (Fig. 1b) using a hand shovel, and then quickly homogenized before the storage in 130 the plastic ziplock bag for the analyses of sediment OC and microbe. At a reservoir site 131 132 adjacent to W1, water and surface sediment samples were also collected in triplicate using a stainless-steel bucket and an Ekman grab sampler, respectively. The collected 133 water and sediment samples were kept frozen in an ice box (-5  $\circ$ C--10  $\circ$ C) and 134 transported to the laboratory for analysis within three days. 135

Dissolved oxygen (DO) in the reservoir was measured using a multi-sensor probe
(YSI 6600, Yellow Springs Instruments, USA), and the DO at each well was measured *in situ* using a DO meter (JPB-607A, Shanghai INESA Scientific Instrument Co. Ltd.,
China) by directly placing the probe in the well. Analysis of dissolved organic carbon

(DOC) in the water was conducted on filtered samples (Whatman GF/F, UK) using a
total organic carbon analyzer (Liqui TOC II, Elementar Inc., Germany). Sediment OC
was determined using a vario MACRO cube elementar (Elementar Inc., Germany).
Fresh sediment was freeze-dried and ground before analysis. Approximately 30 mg of
each sample was weighed in a tin cup and acidified with two drops of 8 % H<sub>3</sub>PO<sub>4</sub> to
remove inorganic carbonates before OC analysis.

### 146 **2.4 Methane flux analysis**

Methane fluxes from the reservoir (eight sampling sites) and island (seventeen sampling 147 148 sites) were analyzed using the static chamber method (Duchemin et al., 1999). The static chamber method has been widely used in the analyses of methane fluxes from 149 air-soil/water interfaces, including the low-to-negative methane flux analyses 150 151 (Veldkamp et al., 2013; Wang et al., 2013). The sampling sites are shown in Fig. 1b and Fig. S4. The plexiglass chamber consisted of a 6.28-L cylinder (20 cm in diameter, 152 20 cm in height) and a removable Styrofoam collar, and was covered with Mylar paper 153 154 to prevent temperature rise inside the chamber when exposed to the sun. During gas collection in the reservoir, the chamber was fitted with the Styrofoam collar, which 155 maintained the upper closed portion of the chamber about 10 cm above the water 156 surface (Fig. S5). The outlet of the chamber was open during the chamber deployment, 157 and was left to stand for 20 min to equilibrate with ambient pressure outside before 158 sample collection. During gas collection on the island, the chamber was carefully 159 inserted 5 cm deep into the sediment according to (Smith et al., 2000), leaving 15 cm 160 above the sediment surface. The outlet of the chamber was also open during the 161

chamber deployment, and was left to stand for 90 min to equilibrate before sample 162 collection. Gas samples (20 ml) were collected every 10 min over a 40-min period using 163 a 25-ml polypropylene syringe and injected into a 12-ml pre-evacuated Exetainer® vial 164 (839 W, Labco, UK) with double wadded septa caps for storage until analysis using a 165 gas chromatograph (7890B, Agilent Technologies, USA). Exetainer® vials have been 166 evaluated for storage of gas samples and they can limit sample loss to insignificant 167 amounts for at least 3 months (Glatzel and Well, 2008; Van Dam et al., 2018). Gas 168 fluxes were calculated using linear regression based on the concentration changes of 169 170 five samples over time. Linear regression correlation coefficients of less than 0.95 were not accepted for further calculations (Duchemin et al., 1999). Methane fluxes at each 171 site were measured in triplicate by placing three individual chambers. Simple spline 172 173 interpolation was used to interpolate the methane emissions from the sampling sites into space in the reservoir and island separately (Immerzeel et al., 2009), and the range 174 of the uncertain was 0.05 mg h<sup>-1</sup> m<sup>-2</sup>. Methane emission areas at eight different 175 176 categories were also calculated in the island.

177 **2.5 Mic** 

### 2.5 Microbial abundance analysis

After being transported to the laboratory, the frozen sediment samples were stored immediately at -80 °C for further molecular analysis. The sediment methanogens and methanotrophs adjacent to each monitoring well across the island (ten sediment samples) were quantified using qPCR. DNA extraction was undertaken using a FastDNA Power-Max Soil DNA Isolation Kit (MP Biomedical, USA) according to the manufacturer's instructions. The qPCR assay was performed using primers targeting methanogenic

archaeal 16S rDNA (primer set, 1106F/1378R) and methanotrophic pmoA genes 184 (primer set, A189F/M661R) (Watanabe et al., 2007; Ma and Lu, 2011). Gene copies 185 were amplified and quantified in a Bio-Rad cycler equipped with the iQ5 real-time 186 fluorescence detection system and software (version 2.0, Bio-Rad, USA). All reactions 187 were completed in a total volume of 20 µL containing 10 µL SYBR<sup>®</sup> Premix Ex Tag<sup>TM</sup> 188 (Toyobo, Japan), 0.5 mM of each primer, 0.8 μL of BSA (3 mg mL<sup>-1</sup>, Sigma, USA), 189 ddH<sub>2</sub>O, and template DNA. The qPCR program mainly depends on the sequence and 190 length of functional genes and primers used, and different qPCR programs for archaeal 191 192 16S rDNA and *pomA* were applied in this study. The qPCR program for archaeal 16S rDNA was as follows: 95  $^{\circ}$ C for 60 s, followed by 40 cycles of 95  $^{\circ}$ C for 25 s, 57  $^{\circ}$ C 193 for 30 s, and 72 °C for 60 s, and the qPCR program for *pomA* referred to: 95 °C for 60 194 195 s, followed by 40 cycles of 95 °C for 25 s, 53 °C for 30 s, and 72 °C for 60 s. A standard curve was established by serial dilution (10<sup>-2</sup>-10<sup>-8</sup>) of known concentration plasmid 196 DNA with the target fragment. All PCRs were run in triplicate on 96-well plates (Bio-197 198 Rad, USA) sealed with optical-quality sealing tape (Bio-Rad, USA). Three negative controls without the DNA template were included for each PCR run. 199

200 **2.6 Data analysis** 

One-way analysis of variance (ANOVA) was employed to test the statistical significance of differences between sampling sites. Post-hoc multiple comparisons of treatment means were performed using the Tukey's least significant difference procedure. All statistical calculations were performed using the SPSS (v22.0) statistical package for personal computers. The level of significance was P < 0.05 for all tests.

## 206 **3 Results**

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## 207 **3.1 Physicochemical characteristics**

209 with that of the bulk reservoir water. Lateral gradients of groundwater DO and DOC

As shown in Fig. 2, the island groundwater had lower DO but higher DOC, compared

210 were observed in the island. From the island edge to the center, DO and DOC decreased

significantly from 50.7  $\pm$  2.0 to 4.5  $\pm$  0.9% and 7.30  $\pm$  0.54 to 1.70  $\pm$  0.39 mg L<sup>-1</sup>,

respectively (P < 0.05) (Fig. 2a,b). In general, sediment OC was higher near the island

edge, decreasing from 6.37  $\pm 0.69$  mg g<sup>-1</sup> at the edge to 2.42  $\pm 0.60$  mg g<sup>-1</sup> at the center

of the island. Sediment OC in the reservoir was 6.63  $\pm 0.09$  mg g<sup>-1</sup> (Fig. 2c).

# **3.2 Water level fluctuation and hyporheic exchange**

The reservoir stage fluctuated frequently during the field survey, showing three distinct 216 217 peaks, with a maximum of 3.80 m in the first 37 h and gradual decline to below 1.30 m in the next 60 h, yielding a maximum oscillation of 2.54 m. Similar oscillations were 218 observed in the island water table, but were damped and lagged relatively to the 219 220 reservoir stage fluctuations (Fig. 3a). In W5, W7, and W10, the water levels reached 3.27, 3.41, and 3.33 m, then fell to 1.74, 2.09, and 2.01 m, for a maximum oscillation 221 of 1.53, 1.33, and 1.32 m, respectively. Data from the automated water level recorders 222 indicated that the water level responses in W5, W7, and W10 lagged the reservoir stage 223 by 20, 25, and 30 min, respectively. Lateral hyporheic exchanges across the island bank 224 were calculated according to the Darcy Law, showing that the flux was largest at the 225 226 island edge and decreased from the edge to the center. The water exchange across the 0-10.5 m island edge zone was 1.2 and 4.7 times higher than those across the 10.5-20.5 227

m and 20.5–35.5 m zones, respectively. The flow rates at the reservoir-W5, W5-W7,

and W7-W10 zones were relatively consistent at -0.55–1.35, -0.89–0.28, and -0.39–

230  $0.17 \text{ m}^2 \text{ d}^{-1}$  (Fig. 3b), resulting in a water exchange volume of 2.61, 2.26, and 0.56 m<sup>3</sup>,

- respectively, over the 115-h observation period.
- 232 **3.3 Methane emissions**

High methane emission rates were observed at the island center, with a maximum of 233 10.4 mg h<sup>-1</sup>m<sup>-2</sup>. However, a ring-like zone of low-to-negative methane emission 234 appeared at the drawdown area around the island edge, where the methane flux was 235 maintained at  $-0.2-1.6 \text{ mg h}^{-1}\text{m}^{-2}$  (Fig. 4a). The negative flux values also suggest the 236 occurrence of a methane sink at the island edge. The ring-like zone accounted for 89.1 % 237 of the island area, of which 9.1 % accounted for the methane sink zone (Fig. 4b). 238 Methane emissions were estimated to 8.3 and 5.4 g  $h^{-1}$  in total from the island center 239 and the ring-like zone, accounting for 60.6% and 39.4%, respectively. Compared with 240 the island, the methane flux from the adjacent reservoir was moderate at  $0.4-5.5 \text{ mg h}^{-1}$ 241  $^{1}m^{-2}$  (Fig. 4a). 242

# 243 **3.4 Methanogen and methanotroph abundances**

Methanogens and methanotrophs were distributed non-uniformly across the island. In general, methanogen counts were low at the island edge but high at the center, whereas methanotrophs were abundant at the island edge but scarce in the center. From the island edge to the center, the methanogenic archaeal 16S rDNA gene increased from  $0.12 \times$  $10^5$  to  $5.34 \times 10^5$  copies g<sup>-1</sup>, and the methanotrophic *pmoA* gene decreased from  $1.57 \times$  $10^6$  to  $0.64 \times 10^6$  copies g<sup>-1</sup> (Fig. 5a). The ratio of methanogen to methanotroph abundance increased from 0.01 at the island edge to 0.83 at the center (Fig. 5b).

251

# 252 **4 Discussion**

## **4.1 Hyporheic exchange and oxygen gradients**

In hydropower reservoirs, the release of water pulses is often employed to increase 254 power production and meet daily electricity peak demand (Bonalumi et al., 2012; 255 Toffolon et al., 2010). Such hydropeaking creates daily water level fluctuations in the 256 reservoir. In this study, frequent water level fluctuations were observed within the 115-257 258 h observation period, with a maximum of 3.80 m (Fig. 3a). In presence of head gradients and sediment resistance (Gerecht et al., 2011), a hysteretic response occurred in the 259 island bank water table (Fig. 3a), driving water exchange between the reservoir and 260 261 island (Fig. 3b). The water exchange flux was largest close to the island edge and decreased from the edge to the center, as water table fluctuations were attenuated (Fig. 262 3a). 263

264 During a storage-release cycle, the island switched from water gaining to losing at daily or hourly scales, creating a ring-like drawdown area of enhanced hyporheic 265 exchange around the island. The drawdown area extended tens of meters into the island 266 bank (Fig. 3b). If the river system was unregulated, however, hydrodynamics within the 267 drawdown area would likely exhibit seasonal or annual patterns, or keep pace with 268 snowmelt and rainstorm events, under a natural base flow-fed regime. In this case, the 269 270 drawdown area may be limited or altogether absent (Boano et al., 2008; Cardenas and Wilson, 2007). 271

Exchange across the sediment-water interface involves mixing of surface water and 272 groundwater through hyporheic flow (Hester et al., 2013; Naranjo et al., 2015). In this 273 274 study, when the reservoir water entered the hyporheic flow path, it was typically rich in oxygen (Fig. 2d). As oxygen was consumed through aerobic respiration, other terminal 275 electron acceptors were utilized (Klupfel et al., 2014), creating an oxygen gradient 276 along the hyporheic flow path (Fig. 2d). This caused the spatial heterogeneity of the 277 abundances of oxygen-sensitive methanogens and methanotrophs across the island. The 278 oxygen-rich environment at the island edge favored methanotroph growth and inhibited 279 280 methanogen growth, while the oxygen-poor environment at the island center inhibited methanotroph growth and favored methanogen growth. As a result, we detected low 281 methanogen abundance at the island edge, but high abundance at the center, with 282 283 methanotrophs showing the opposite pattern (Fig. 5).

## **4.2 Spatial heterogeneity of methane emissions**

In dammed rivers, riverbed sediment accumulation creates potential methane emission 285 286 hotspots. In this study, however, high methane emissions were only observed at the 287 island center, with a ring-like low methane emission zone or even methane sink appearing around the island edge (Fig. 4a). This was attributed to the spatial 288 heterogeneity of methanogens and methanotrophs across the island (Fig. 5), leading to 289 an increase in methane production and a decrease in methane consumption from the 290 island edge to the center. The methane sink at the island edge (Fig. 4) was mainly 291 292 attributed to the strong oxidation by methanotrophs, which may consume methane to below equilibrium with the atmosphere. Methane emissions may not only rely on 293

bacterial abundance but also bacterial activity (Schwarz et al., 2008). This deserves further studies using other molecular biology techniques, such as DNA/RNA-based stable isotope probing (Dumont and Murrell, 2005). In addition, groundwater DOC and sediment OC at the island edge, which are carbon sources for methane emission, were higher than that at the island center (Fig. 2b,c), suggesting that both sediment heterogeneity and dilution effects of hyporheic exchange had limited contribution to the spatial pattern of methane emissions in the island.

#### 301 **4.3 Implications**

302 Greenhouse gas emissions significantly detract from the green credentials of hydropower, and have thus received considerable research attention (Giles, 2006; Hu 303 and Cheng, 2013). Previous studies have revealed that damming causes significant 304 305 retention of carbon and creates deep, anoxic sediment strata, fueling methanogenesis and net water-air methane flux (Maeck et al., 2013). This study demonstrated that 306 methane emissions at the most area of the sediment-deposited island were generally 307 308 lower than the adjacent reservoir under water level fluctuation induced by reservoir operation (Fig. 4a), but higher than the drawdown area at other reservoir bank (-0.08– 309  $0.66 \text{ mg h}^{-1}\text{m}^{-2}$ ), such as Three Gorge Reservoir (Chen et al., 2011). This was mainly 310 due to the deep sediment strata (about 60 m in depth) in the island. Given the widely 311 distributed sediment-deposited islands in reservoirs, it should be of concern in future 312 estimations of greenhouse gas emissions from dammed rivers. 313

Until now, few studies have concentrated on organic carbon mineralization in the drawdown area in reservoirs, with most focusing on the process of denitrification

(Zarnetske et al., 2011b). Carbon emissions in the drawdown area are poorly understood, especially in regulated and dammed rivers. This study fills the knowledge gap and adds to our understanding of the ecological impacts of hydropower exploitation. Under water level fluctuation induced by reservoir operation, variable oxygen conditions and methane production may also affect the mercury cycle in the drawdown area and thereby the release of methylmercury (a bioaccumulative environmental toxicant) to the river (Marvin-DiPasquale et al., 2009), a subject deserving of further study.

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### 324 **5** Conclusions

In dammed rivers, sediment deposited islands are widely distributed in sidebays and are 325 potential hotspots of methane emission to the atmosphere. In this study, high methane 326 327 fluxes were only observed at the island center, while a ring-like zone of low methane emission or even sink was found in the drawdown area around the island edge. We 328 attribute this spatial heterogeneity of methane emissions to hyporheic exchange 329 330 between the reservoir and island. Under reservoir operation, frequent water level fluctuations drove hyporheic exchange, creating oxygen gradients along the hyporheic 331 flowpath. These oxygen gradients affected the microbial communities associated with 332 methane production and consumption, producing the spatial heterogeneity in methane 333 emissions across sediment-deposited islands. This study will help us to evaluate the 334 global warming effects of hydropower systems. 335

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338	Data	availa	bility

339 The data presented here can be obtained upon request to Wenqing Shi (wqshi@nhri.cn).340

341	Author	contri	bution

- 342 Qiuwen Chen designed the research; Wenqing Shi, Yuyu Ji and Yuchen Chen performed
- 343 the research; Cheng Chen and Juhua Yu contributed new reagents/analytic tools;
- 344 Qiuwen Chen and Wenqing Shi analyzed the data; Jianyun Zhang and Bryce R. Van
- 345 Dam contributed significant discussions and inputs; Wenqing Shi and Qiuwen Chen
- 346 wrote the paper with input from all authors.

# 347 **Competing interests**

- 348 The authors declare that they have no conflict of interest.
- 349

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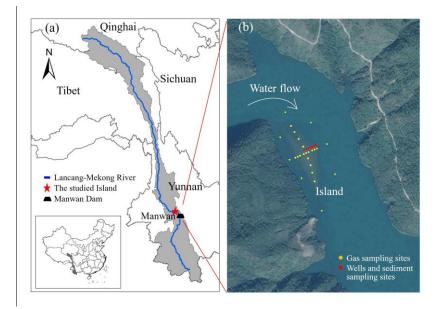
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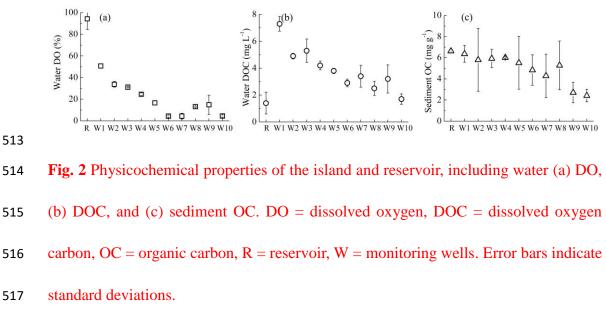


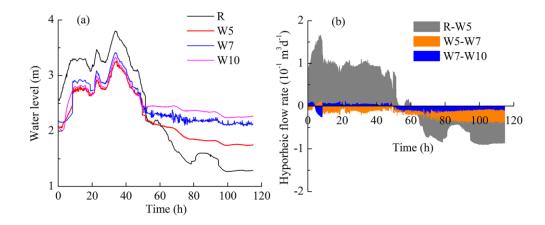


**Fig. 1** Map of the studied island in Manwan reservoir, Lancang-Mekong River. (a) The

511 island site in the river; (b) The monitoring wells, sediment and gas sampling sites on

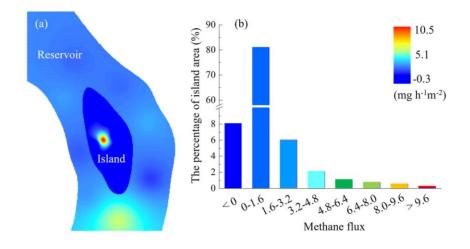
512 the island.





**Fig. 3** Vertical water level fluctuation (a) and horizontal hyporheic flow rate (b). R =reservoir, W = monitoring wells. Positive fluxes indicate net flow from the reservoir to island, whereas negative values indicate net flow from the island to reservoir.

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525 Fig. 4 Methane emissions from the island and reservoir. (a) Spatial pattern of methane

emissions; (b) Percentage of the island area emitting methane at a certain flux. Methane

527 fluxes were interpolated separately for the island and reservoir.

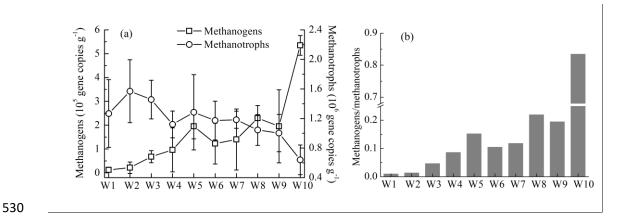


Fig. 5 Abundances of sediment methanogens and methanotrophs in the island. (a)
Spatial patterns of methanogen and methanotroph abundances across the island; (b) The
ratio of methanogen to methanotroph abundance at each site. W= monitoring wells.
Error bars indicate standard deviations.



**Fig. S1** The river impoundment formed islands in the Lancang-Mekong River. There were about 33 sediment-deposited islands, of which 81.8% are located at the convex bank (a), and 18.2% at the concave bank (b).

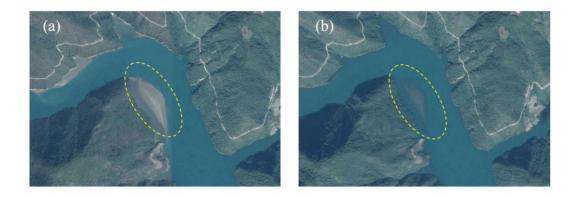


Fig. S2 Island image before (a) and after (b) flooding under reservoir operation.

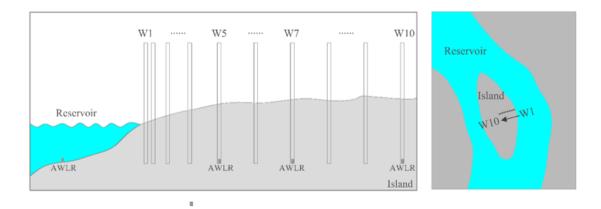


Fig. S3 Monitoring wells established from the island edge to the center. Sediment samples were collected adjacent to each well.

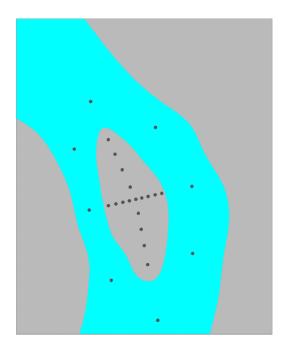
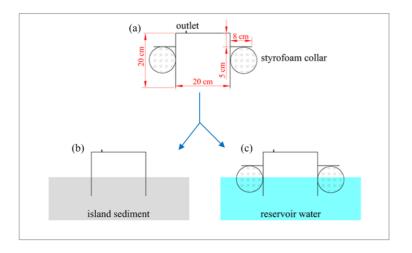


Fig. S4 Sampling sites for methane flux analyses in the island and reservoir, including eight sites in the reservoir and seventeen sites on the island.



**Fig. S5** Bifunctional static chambers (a) for methane flux analyses across the sediment-air interface in the island (b) and water-air interface in the reservoir (c). Styrofoam collar

was removable.

Sites		The con	centration	in the char	nber (×10 <sup>-6</sup>	<sup>6</sup> mol/L)	Flux	Flux	Flux	Average flux
		0 min	10 min	20 min	30 min	40 min	(×10 <sup>-6</sup> mol/h)	(×10 <sup>-6</sup> mol/h m <sup>2</sup> )	( <b>mg/h m</b> <sup>2</sup> )	( <b>mg/h m</b> <sup>2</sup> )
R1	Rep-1	0.452	0.773	1.174	1.375	1.656	1.806	57.5	0.92	1.07
	Rep-2	0.641	0.921	1.280	1.680	2.160	2.279	72.6	1.16	
	Rep-3	0.541	1.106	1.272	1.637	2.103	2.193	69.8	1.12	
	Rep-1	0.797	0.887	1.232	1.778	2.081	2.075	66.1	1.06	1.13
R2	Rep-2	1.166	1.431	1.722	2.112	2.778	2.343	74.6	1.19	
	Rep-3	0.823	0.869	1.197	3.825	4.273	NA	NA	NA	
	Rep-1	0.678	1.075	1.809	2.243	3.264	3.804	121.1	1.94	
<b>R3</b>	Rep-2	0.822	1.261	2.282	2.702	3.704	4.323	137.7	2.20	2.10
	Rep-3	1.243	1.543	2.246	2.969	4.043	4.216	134.3	2.15	
	Rep-1	0.573	0.660	0.759	1.018	1.189	0.954	30.4	0.49	
<b>R4</b>	Rep-2	0.390	0.553	0.665	0.797	0.829	0.673	21.4	0.34	0.39
	Rep-3	0.476	0.622	0.762	0.863	0.909	0.665	21.2	0.34	
	Rep-1	1.867	2.871	4.802	6.432	8.739	10.383	330.7	5.29	
R5	Rep-2	1.294	2.554	4.565	6.582	9.349	12.082	384.8	6.16	5.47
	Rep-3	1.188	2.165	4.289	4.913	7.935	9.745	310.4	4.97	
	Rep-1	0.493	1.175	1.608	2.142	2.678	3.203	102.0	1.63	
R6	Rep-2	0.881	1.411	1.881	2.751	3.561	4.021	128.1	2.05	1.84
	Rep-3	0.677	1.432	1.381	2.988	3.463	NA	NA	NA	
	Rep-1	0.388	0.802	0.923	1.224	1.482	1.566	49.9	0.80	0.84
<b>R7</b>	Rep-2	0.653	0.876	1.875	1.896	2.165	NA	NA	NA	
	Rep-3	0.794	0.930	1.420	1.609	1.902	1.737	55.3	0.89	
	Rep-1	0.971	1.626	2.515	3.401	4.015	4.718	150.3	2.40	1.84
<b>R8</b>	Rep-2	0.879	0.997	1.377	1.995	2.469	2.508	79.9	1.28	
	Rep-3	0.726	0.726	1.543	1.945	2.807	3.605	114.8	1.84	
	Rep-1	0.463	0.434	0.358	0.295	0.200	-0.399	-12.7	-0.20	
1	Rep-2	0.454	0.414	0.358	0.255	0.157	-0.452	-14.4	-0.23	-0.21
	Rep-3	0.386	0.314	0.258	0.214	0.139	-0.356	-11.3	-0.18	
	Rep-1	0.878	1.213	1.556	2.009	2.810	2.796	89.0	1.42	1.26
2	Rep-2	0.784	0.970	1.432	1.945	2.442	2.575	82.0	1.31	
	Rep-3	0.993	1.187	1.534	1.898	2.358	2.065	65.8	1.05	
	Rep-1	1.068	1.483	2.425	3.567	4.738	5.654	180.1	2.88	2.78
3	Rep-2	1.282	1.919	2.799	3.304	4.482	4.671	148.8	2.38	
	Rep-3	1.482	1.949	2.876	3.967	5.519	6.055	192.8	3.09	
-	Rep-1	2.893	6.897	8.251	12.608	16.811	20.13	641.1	10.26	10.41
4	Rep-2	3.892	7.996	10.451	15.906	17.211	20.73	660.2	10.56	
	Rep-3	1.278	2.301	4.061	5.868	8.291	NA	NA	NA	
	Rep-1	0.786	NA	2.784	4.324	NA	NA	NA	NA	
5	Rep-2	0.876	1.805	3.180	4.625	5.769	7.564	240.9	3.85	4.27
	Rep-3	1.069	2.075	3.208	5.141	7.203	9.201	293.0	4.69	
6	Rep-1	1.221	1.519	1.697	1.875	1.933	1.068	34.0	0.54	0.49

Table S1 The data about methane fluxes at each sampling site

	Rep-2	0.782	0.819	0.999	1.153	1.338	0.868	27.6	0.44	
	Rep-3	1.887	1.267	1.756	2.154	2.375	NA	NA	NA	
	Rep-1	0.456	0.476	0.481	0.493	0.507	0.071	2.3	0.04	
7	Rep-2	0.479	0.489	0.499	0.505	0.515	0.05	1.6	0.03	0.03
	Rep-3	0.578	0.587	0.602	0.611	0.62	0.065	2.1	0.03	
	Rep-1	0.399	0.408	0.417	0.422	0.428	0.043	1.4	0.02	
8	Rep-2	0.456	0.471	0.482	0.503	0.512	0.086	2.7	0.04	0.03
	Rep-3	0.488	0.497	0.505	0.517	0.519	0.049	1.6	0.02	
	Rep-1	0.678	0.6768	0.676	0.675	0.6745	-0.005	-0.2	0.00	
9	Rep-2	0.544	0.536	0.531	0.528	0.521	-0.032	-1.0	-0.02	-0.01
	Rep-3	0.487	0.483	0.48	0.475	0.472	-0.023	-0.7	-0.01	
	Rep-1	0.661	0.67	0.678	0.683	0.687	0.039	1.2	0.02	
10	Rep-2	0.495	0.502	0.506	0.512	0.519	0.035	1.1	0.02	0.02
	Rep-3	0.785	0.791	0.795	0.802	0.809	0.0355	1.1	0.02	
	Rep-1	0.555	0.5545	0.5538	0.5535	0.5528	-0.003	-0.1	0.00	
11	Rep-2	0.453	0.454	0.454	0.455	0.456	0.004	0.1	0.00	0.00
	Rep-3	0.768	0.745	0.755	0.777	0.815	NA	NA	NA	
	Rep-1	0.787	1.199	1.443	2.108	2.588	2.707	86.2	1.38	
12	Rep-2	0.722	0.805	1.119	1.403	1.844	1.705	54.3	0.87	1.12
	Rep-3	0.477	0.674	0.846	0.876	1.532	NA	NA	NA	
	Rep-1	0.491	1.118	1.694	2.171	2.847	3.46	110.2	1.76	
13	Rep-2	0.493	1.015	1.290	1.965	2.390	2.85	90.8	1.45	1.61
	Rep-3	NA	0.564	NA	0.762	1.438	NA	NA	NA	
	Rep-1	0.393	0.444	0.477	0.521	0.532	0.021	0.7	0.01	
14	Rep-2	0.420	0.425	0.430	NA	0.439	0.029	0.9	0.01	0.01
	Rep-3	0.422	0.427	0.432	0.435	0.440	0.0265	0.8	0.01	
	Rep-1	0.754	0.733	0.695	0.677	0.644	-0.166	-5.3	-0.08	
15	Rep-2	0.472	0.463	0.455	0.447	0.435	-0.054	-1.7	-0.03	-0.05
	Rep-3	0.511	0.493	0.475	0.457	0.447	-0.098	-3.1	-0.05	
	Rep-1	0.567	0.540	0.525	0.511	0.503	-0.094	-3.0	-0.05	
16	Rep-2	0.853	0.840	0.835	0.821	0.809	-0.064	-2.0	-0.03	-0.04
	Rep-3	0.454	0.440	0.435	0.411	0.398	-0.085	-2.7	-0.04	
	Rep-1	0.263	0.564	0.672	0.675	0.785	NA	NA	NA	
17	Rep-2	0.567	0.570	0.572	0.574	0.575	0.012	0.4	0.01	0.01
	Rep-3	0.368	0.370	0.375	0.377	0.381	0.02	0.6	0.01	