Changes in Methane Emission and Community Composition of Methane-Cycling Microorganisms Along an Elevation Gradient in the Dongting Lake Floodplain, China

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Abstract: Methane (CH$_4$) emission and environmental controls of CH$_4$-cycling microorganisms are unclear in inland floodplains. Here, we examined soil CH$_4$ emissions and the community composition of CH$_4$-cycling microorganisms under three vegetation types—mudflat (MF, no vegetation cover), Carex meadow (CM, mainly Carex brevicuspis), and reed land (RL, mainly Miscanthus sacchariflorus)—from water-adjacent areas to higher-elevation land in the Dongting Lake floodplain, China. The results showed that CH$_4$ emission is the highest in CM, while significant absorption was observed in the RL site. The abundance ratio of methanogen/methanotroph was the highest in CM, intermediate in MF, and lowest in RL. The Methanosarcinaceae family represented the dominant methanogens in the three sampling sites (41.32–75.25%). The genus Methylocystis (60.85%, type II methanotrophs) was dominant in CM, while Methylobacter and Methylosarcina (type I methanotrophs) were the dominant genera in MF (51.00%) and RL (50.24%), respectively. Structural equation model analysis showed that methanogen and methanotroph abundance were affected by water table depth, soil water content, and pH indirectly through soil organic content, total nitrogen, microbial biomass carbon, and microbial biomass nitrogen. These results indicated that the Dongting Lake floodplain may change from a CH$_4$ source to a CH$_4$ sink with vegetation succession with an increase in elevation, and the methanogen/methanotroph ratio can be used as a proxy for CH$_4$ emission in wetland soils. The continuous increase in reed area combined with the decrease in Carex meadow may mitigate CH$_4$ emission and enhance the CH$_4$ sink function during the non-flood season in the Dongting Lake floodplain.

Keywords: methanotroph; methanogen; Dongting Lake floodplain; vegetation type; methane flux
1. Introduction

Methane (CH$_4$) is an important greenhouse gas with a global warming potential 28 times higher than that of carbon dioxide (CO$_2$) on a 100-year scale [1]. Natural wetlands are considered the main natural CH$_4$ source for the atmosphere because of their high productivity and low redox capacity. The annual CH$_4$ emission from natural wetlands is reported to be 172 Tg CH$_4$•y$^{-1}$ (1σ SD, ± 12 Tg CH$_4$•y$^{-1}$), accounting for approximately 62% of the annual global (natural) CH$_4$ budget [2,3]. Although CH$_4$ emissions and microbial production mechanisms from high-latitude peatlands have been well studied [4–8], relatively little data have been published on the low-latitude wetlands, which is the most important source of uncertainty on the methane budget [2].

Low-latitude seasonal floodplains are prolific sources of CH$_4$ because of their substantial net primary productivity and high seasonal temperatures [9]. For example, the large CH$_4$ emission in the Amazon floodplain trees adapted to permanent or seasonal inundation can account for about 15% of the global wetland CH$_4$ emission [10]. Water regime, i.e., water table depth (WTD), soil water content (SWC), etc., and C and N concentrations in the soil, i.e., soil organic content (SOC), nitrogen, dissolved organic carbon (DOC), etc., along with the elevation gradient, may act as critical factors influencing CH$_4$ production and emission in floodplains [11–13]. It has been documented that the water regime was a predominant factor affecting the methanogenic archaea (methanogens) and CH$_4$-oxidizing microbes (methanotrophs) community over regional and seasonal variations [14]. Plant zonation along an elevation is a common phenomenon in floodplain wetlands. Vegetation type can also play a critical role in determining the community structures of methanogens and methanotrophs because it can provide sufficient substrates (i.e., soil C, N, etc.) for the growth of CH$_4$-cycling microorganisms by plant litter decomposition and root exudate [15,16]. Nevertheless, with an increase in elevation, methanogen should be unlikely to survive owing to the WTD and SWC decline and increase in oxygen availability [17]. Therefore, the effects of vegetation and water regime, along with an elevation gradient, on CH$_4$-cycling microorganisms, are still conflicting.

Currently, under the double pressure of anthropogenic disturbances and global climate change, the vegetation distribution pattern in wetlands changes significantly owing to variation in the water regime, especially in river-connected floodplains [18,19]. For example, in the East Dongting Lake wetlands, the area covered by reed and forest increased significantly, but that of Carex meadow (CM) showed a sharply decreasing trend during 1995–2015, particularly after the implementation of the Three Gorges Dam (TGD) upstream of the Yangtze River [19]. In this study, we analyzed the CH$_4$ flux and the community structure of methanogens and methanotrophs along an elevation gradient in the Dongting Lake floodplain. Three vegetation types—mudflat (MF, no vegetation cover), CM (dominated by Carex brevicuspis), and reed land (RL, dominated by Miscanthus sacchariflorus)—were distinctly distributed from water-adjacent areas to higher-elevation land. The objectives of our study were as follows: (1) to analyze the CH$_4$ emission from three vegetation types along an elevation gradient; (2) to elucidate the diversity and community structure of methanogens and methanotrophs in three vegetation types and vertical soil profiles; and (3) to quantify the relationships between environmental variables and abundance of methanogens and methanotrophs in three vegetation types.

2. Experiments

2.1. Study Site

Dongting Lake (28°30′–30°20′ N, 111°40′–113°10′ E), with an area of 2625 km$^2$, is the second-largest inland freshwater lake in China. It is connected to the Yangtze River by distributary channels (Songzi, Taiping, and Ouchi) and has four other water inlets (Rivers Xiang, Zi, Yuan, and Li) in the Hunan Province (Figure 1). The wetlands of this lake are characterized by large seasonal water level fluctuations (12–14 m), with the maxima in August and minima typically in January or February, which provide the basic hydrological regime for maintaining large areas of floodplain [20]. The floodplains are completely flooded from June to October and are exposed from November to the following May [21].
The study was conducted at a monitoring sampling site (29°30′ N, 112°48′ E) of Dongting Lake Station for Wetland Ecosystem Research, a member of the Chinese Ecosystem Research Network (CERN). The monitoring site has an area of 200 ha, and it is covered by mudflats (no vegetation cover), Carex communities (mainly Carex brevicuspis), and reed (mainly Miscanthus sacchariflorus), along with a small elevation gradient (22–26 m, Yellow Sea Datum 1985). The mean annual temperature in the study area is 16.4–17.0 °C, with the lowest temperatures in January (3.9–4.5 °C) and the highest in July (28.6–29.1 °C). The annual precipitation is 1382 mm, with more than 60% of rainfall occurring from April to August [22].

Figure 1. Maps of Dongting Lake showing the location and elevation of the sampling area.

2.2. Soil Sampling Process

Soil samples were collected from three sampling sites—MF, CM, and RL—on 10 November 2016, after the flood season. Five sampling plots at intervals of 100 m in a parallel line were randomly selected at each site. Five 30-cm-deep soil cores at each sampling plot were collected using steel sediment samplers (d = 10 cm) and split into three depths: 0–10, 10–20, and 20–30 cm. Each collected soil sample (ca. 1000 g in fresh weight) was divided into two sub-samples. One was stored at 4 °C for physicochemical analysis (ca. 900 g in fresh weight), and the other was stored in liquid nitrogen for DNA extraction and molecular analysis (ca. 100 g in fresh weight).

2.3. Soil Physical and Chemical Analysis

Soil pH was determined using a pH meter (Mettler Toledo, Shanghai, China) with a soil-to-water ratio of 1:2.5. The SWC was determined by the mass ratio of the pore water to solids by drying the samples in the oven at 105 °C for 48 h [23]. The SOC was determined by wet oxidation with KCr2O7 + H2SO4 and titration with FeSO4. Ammonium (NH4+) and nitrate (NO3−) were determined by the Flow Injection Analyzer AA3 (SEAL, 22844 Norderstedt, Germany) after extraction with 2M K2SO4 solution. Total carbon (TC) and total nitrogen (TN) were measured by the Element Analyzer (Elementar, Munich, Germany). Soil DOC was measured as follows: each 10-g wet-soil sample was incubated with 50 mL of distilled water for 30 min on an end-over-end shaker and then centrifuged for 20 min at 1680× g (5000 rpm). The extracted solutions were passed through a 0.45-μm filter membrane and analyzed using a total organic carbon (TOC) Analyzer (TOC-VWP, Shimadzu, Kyoto, Japan) [24]. The estimation of soil microbial biomass carbon (MBC) and microbial biomass nitrogen (MBN) was performed by the chloroform–fumigation–extraction method: 25 g of fresh soil was
fumigated, and carbon dissolved in 0.5 M K$_2$SO$_4$ extracts was measured by automated analyses on a TOC Analyzer (TOC-VWP, Shimadzu, Kyoto, Japan) [25].

2.4. Gas Sampling and CH$_4$ Lux Calculation

The gas collection was carried out at the three sampling sites every 7 days from 5 November to 6 December 2016, using the static opaque chamber–gas chromatography method [26]. The chambers (40 cm in diameter and 50 cm in height) were made of 0.5-cm-thick opaque Polyvinyl chloride (PVC) pipe, with a volume and area of 0.0628 m$^3$ and 0.1256 m$^2$, respectively. Because the reeds were harvested for paper making during the non-flooding season of the Dongting Lake floodplain, the CH$_4$ emission in RL during our sampling period only represents the emission in the soil and belowground plant tissues. The chambers were equipped with a small fan for gentle air circulation and wrapped with thermal insulating Styrofoam and radiation-reflecting tinfoil to prevent dramatic temperature changes in the headspace during chamber closure. A PVC chamber base (height = 10 cm; diameter = 41 cm) was permanently inserted into the soil at each spatial replicate.

On the day of observation, gas sampling at the three sites was simultaneously performed between 8:00 and 10:00 a.m., when the gas flux and air temperature were representatives of the daily averages. At each sampling time, the chamber was sealed, and five gas samples (30 mL each) were collected using a polypropylene syringe at 10-min intervals. The samples were injected into labeled evacuated Exetainers (Labco Ltd., London, UK). The CH$_4$ concentrations were analyzed with a gas chromatograph (Agilent 7890A, Santa Clara CA, USA) equipped with a flame ionization detector (FID). The gas fluxes were calculated by the following formula:

$$F_{CH_4} = \frac{dc}{dt} \cdot 0.53 \cdot \frac{273}{T} \cdot \frac{P}{1013} \cdot \frac{H}{1000}$$

where $F_{CH_4}$ is flux (µg C m$^{-2}$ h$^{-1}$), and $dc/dt$ is the slope of the linear regression of gas concentration at time approaching zero. T is the absolute temperature (K) during sampling. P is the atmospheric pressure (KPa) during the sampling period. H is the height (m) of the chamber headspace.

2.5. DNA Extraction, PCR Amplification, and High-Throughput Sequencing

DNA from soil samples was extracted according to the manufacturer’s instructions using the FastDNA® Spin Kit for Soil (MP Biomedicals, Irvine, CA, USA). The extracted DNA was evaluated by electrophoresis on a 1% agarose gel and quantified by spectrophotometry (NanoDrop, Waltham, MA, USA).

A 470-bp mcrA gene fragment, which generally represents methanogens community, was PCR-amplified from community DNA samples by means of a forward primer MLfF (5′-GGTGGTGTTGACGACTCAGCACGAC-3′) and reverse primer MLrR (5′-TTCTTATTGGCAGATTTGACTAG-3′) [27]. A 472-bp pmoA gene fragment, which generally represents methanotrophs community, was PCR-amplified from community DNA samples by means of a forward primer A189F (5′-GGNGACTGGGACTTCTGG-3′) [28] and reverse primer mb661R (5′-CCGGGMCGAGCTCTTTGACC-3′) [29]. The mcrA and pmoA gene fragments were PCR-amplified by an ABI GeneAmp® 9700 PCR thermocycler (ABI, Vernon, CA, USA) with an eight-base sequence barcode unique to each sample at the 5′ end of the MLfF/MLrR and A189F/mb661R primer pairs, respectively. Amplification was performed in a 20-µL final volume containing 4 µL of 5 × FastPfu Buffer, 2 µL of 2.5 mM dNTPs, 0.4 µL of FastPfu Polymerase, 0.2 µL of BSA, 0.8 µL of each primer (final concentration 5 µM), and 10 ng of extracted DNA. The thermal cycling conditions were as follows: 95 °C for 3 min followed by 35 cycles (mcrA) and 36 cycles (pmoA) of denaturing at 95 °C for 30 s, annealing at 55 °C for 30 s, and extension at 72 °C for 45 s. Final extension reactions were carried out for 10 min at 72 °C. The PCR products were extracted and further purified using the AxyPrep DNA Gel Extraction Kit (Axygen Biosciences, New York, NY, USA). The resulted PCR products of the same sample were mixed and analyzed by 2% agarose gel electrophoresis. Purified amplicons were pooled
in equimolar concentrations and paired-end sequenced (2 × 300) on an Illumina MiSeq PE300 platform at Majorbio Bio-Pharm Technology Co. Ltd. (Shanghai, China) (Figure S1). The raw reads were deposited into the NCBI Sequence Read Archive (SRA) database (Accession Number: PRJNA497518).

2.6. Real-Time Quantitative PCR

Methanogens and methanotrophs were quantified by ABI 7500 Real-Time PCR software (ABI, California, USA) using the primer pair MLfF/MLrR and A189F/mb661R. The PCR system contained a 2X ChamQ SYBR Color qPCR Master Mix (16.5 µL), the forward primer (5 µM, 0.8 µL), the reverse primer (5 µM, 0.8 µL), template DNA (19.1 ng for mcrA and 21.3 ng for pmoA, 2 µL), and ddH2O (7.4 µL). Quantitative PCR was carried out as follows: 5 min at 95 °C for initial denaturation, 35 cycles of 5 s at 95 °C, 30 s at 53 °C (mcrA) and 58 °C (pmoA), and 40 s at 72 °C. After the PCR procedure, three products were randomly selected to be analyzed on an agarose gel and sequenced to check the specificity of the products. Moreover, the melting-curve analysis for each sample was employed to ensure the specificity of each amplicon. A standard curve for the mcrA and pmoA genes was prepared using a 10-fold dilution series (10^2–10^9 copies) of the plasmids containing the target gene fragments. The efficiencies of qPCR for the mcrA and pmoA genes were 89.27% (R^2 = 0.9999) and 97.24% (R^2 = 0.9999), respectively. All qPCRs were run in triplicate with the DNA extracted from each soil sample.

2.7. Data Analysis

Quality control of the amplicon sequences was performed using QIIME pipeline (version 1.9.0). Briefly, low-quality sequences (<50 bp in length with an average quality score of <20) were excluded from further analyses. After the removal of the barcode and primer sequences, operational taxonomic units (OTUs) were defined with an identity threshold of 97% using Usearch (version 7.0). Taxa were assigned using the RDP fungene database (release 7.3). Subsequently, community bar plots and the plot of average CH\textsubscript{4} flux (n = 5) were constructed using Origin V8.0 (Origin Lab, Northampton, MA, USA). The differences in soil properties (SWC, MBC, MBN, DOC, SOC, pH, NH\textsubscript{4}+, NO\textsubscript{3}−, TC, and TN) and the abundance of methane-cycling microorganisms (pmoA and mcrA gene copies) among the three soil layers and three sampling sites (mean value of the soil properties and methane-cycling microorganisms of the whole soil column) were tested by one-way analysis of variance (ANOVA) using R (version 3.4.3). Redundancy analysis (Koh, #46) with 999 Monte Carlo permutation tests was conducted by R (version 3.4.3). Phylogenetic trees were constructed by FastTree (version 2.1.3) using the approximately-maximum-likelihood method and then drawn by R (version 3.4.3). We further fitted a piecewise structural equation model (SEM) using R (version 3.4.3); the “piecewise SEM” package was used to infer the relative importance of environmental factors (WTD, SWC, and pH), C and N availability in the soil (SOC, TN), MBC, and MBN on the abundance of methanogens and methanotrophs. Compared with the traditional variance–covariance-based SEM, the piecewise SEM could (1) piece multiple separate (generalized) linear models together to a single causal network, (2) use Shipley’s test of d-separation to test whether any paths are missing from the model, and (3) use Akaike information criterion (AIC) to compare nested models and for small sample size [30].

3. Results

3.1. CH\textsubscript{4} Flux and Abundance of Methanogens and Methanotrophs

The average CH\textsubscript{4} flux was significantly higher in CM (57.42 ± 39.17 µg C m\textsuperscript{-2} h\textsuperscript{-1}) than in MF (6.32 ± 4.12 µg C m\textsuperscript{-2} h\textsuperscript{-1}) and RL (–14.80 ± 6.83 µg C m\textsuperscript{-2} h\textsuperscript{-1}) (p < 0.01, Figure 2A).
Table 1. The vertical pattern of the abundance of methanogens (gene copies g\(^{-1}\) dws × 10\(^7\)) and methanotrophs (gene copies g\(^{-1}\) dws × 10\(^5\)) in three sites of the Dongting Lake floodplain.

| Microorganisms | Depth(cm) | MF        | CM         | RL         |
|----------------|-----------|-----------|------------|------------|
| Methanogens    | 0–10 cm   | 0.67 ± 0.34 a | 10.17 ± 1.49 a | 2.29 ± 0.50 a |
|                | 10–20 cm  | 0.06 ± 0.03 b | 4.66 ± 0.45 b  | 1.33 ± 0.72 ab |
|                | 20–30 cm  | 0.31 ± 0.14 ab | 3.61 ± 1.13 b  | 0.41 ± 0.20 b  |
| Average        |           | 0.32 ± 0.11 B | 6.15 ± 1.16 A | 1.43 ± 0.38 B |
| Methanotrophs  | 0–10 cm   | 1.17 ± 0.80 a | 11.73 ± 3.60 a | 3.47 ± 0.76 a  |
|                | 10–20 cm  | 0.45 ± 0.32 a | 1.11 ± 0.50 b  | 1.98 ± 0.78 a  |
|                | 20–30 cm  | 0.18 ± 0.077 a | 0.57 ± 0.20 b  | 1.56 ± 0.78 a  |
| Average        |           | 0.62 ± 0.32 B | 4.47 ± 2.10 A | 2.41 ± 0.48 AB |

Different lowercase letters in the same column within each sampling site indicate significant differences (p < 0.05) among the three soil layers; different uppercase letters between three averages in the same column indicate significant differences (p < 0.05) among the three types of sampling sites. Site abbreviations: MF: mudflat, CM: Carex meadow, RL: reed land.

Figure 2. Average methane flux (A) and the abundance ratio of methanogens and methanotrophs (B) from 5 November to 6 December 2016, at three sites in the Dongting floodplain. The data used in (B) is the average ratio of methanogens and methanotrophs gene abundance across the 0–30 cm soil profile. Site abbreviations: MF: mudflat, CM: Carex meadow, RL: reed land. Different lowercase letters in the same column within each sampling site indicate significant differences (p < 0.05) among the three soil layers.

For methanogens, the average abundance value in CM was (6.15 ± 1.16) × 10\(^7\) gene copies g\(^{-1}\) dws, which was significantly higher than in RL (1.43 ± 0.38) × 10\(^7\) gene copies g\(^{-1}\) dws and MF (0.32 ± 0.11) × 10\(^7\) gene copies g\(^{-1}\) dws (p < 0.05, Table 1). For methanotrophs, the average abundance values in CM and RL ((2.41 – 4.47) × 10\(^5\) gene copies g\(^{-1}\) dws) were significantly higher than that in MF (0.62 ± 0.32) × 10\(^5\) gene copies g\(^{-1}\) dws) (p < 0.05, Table 1). However, the difference in the abundance value of methanotrophs between CM and RL was insignificant (p > 0.05, Table 1). The abundance of methanogens and methanotrophs was the highest in soil depths of 0–10 cm, and both decreased with increase in soil depth. The abundance ratio of methanogens and methanotrophs was significantly higher in CM (481.96 ± 134.33) than in MF (112.81 ± 36.50) and RL (52.60 ± 8.78) (p < 0.01, Figure 2B).
Table 1. The vertical pattern of the abundance of methanogens (gene copies g$^{-1}$dws $\times 10^7$) and methanotrophs (gene copies g$^{-1}$dws $\times 10^5$) in three sites of the Dongting Lake floodplain.

| Microorganisms | Depth(cm) | MF | CM | RL |
|----------------|-----------|----|----|----|
| methanogens    |           |    |    |    |
| 0–10 cm        | 0.67 ± 0.34 a | 10.17 ± 1.49 a | 2.29 ± 0.50 a |
| 10–20 cm       | 0.06 ± 0.03 b | 4.66 ± 0.45 b | 1.33 ± 0.72 ab |
| 20–30 cm       | 0.31 ± 0.14 ab | 3.61 ± 1.13 b | 0.41 ± 0.20 b |
| Average        | 0.32 ± 0.11 B | 6.15 ± 1.16 A | 1.43 ± 0.38 B |

| methanotrophs  |           |    |    |    |
| 0–10 cm        | 1.17 ± 0.80 a | 11.73 ± 3.60 a | 3.47 ± 0.76 a |
| 10–20 cm       | 0.45 ± 0.32 a | 1.11 ± 0.50 b | 1.98 ± 0.78 a |
| 20–30 cm       | 0.18 ± 0.077 a | 0.57 ± 0.20 b | 1.56 ± 0.78 a |
| Average        | 0.62 ± 0.32 B | 4.47 ± 2.10 A | 2.41 ± 0.48 AB |

Different lowercase letters in the same column within each sampling site indicate significant differences ($p < 0.05$) among the three soil layers; different uppercase letters between three averages in the same column indicate significant differences ($p < 0.05$) among the three types of sampling sites. Site abbreviations: MF: mudflat, CM: Carex meadow, RL: reed land.

3.2. Diversity and Richness of Methanogens and Methanotrophs

After the amplification of mcrA genes (methanogen) and pmoA genes (methanotroph), the samples were sequenced by high-throughput sequencing. Among the three sampling sites, the average retrieved sequence amount of the mcrA genes and pmoA genes is 15422–16483 and 13368–14773, respectively. The average read lengths were 440 bp (ranging from 421 to 460 bp) and 488 bp (ranging from 481 to 500 bp) for the mcrA and pmoA genes, respectively (Table S1). The coverage index was higher than 98%, indicating that the sequencing effort encompassed a significant amount of biodiversity in the examined samples. For methanogens, the OTU number was the highest in MF (446–468) and lowest in RL (170–214). The ACE and Chao1, which represented methanogen richness, were highest in MF and lowest in RL. The Shannon diversity index was also highest in MF (4.68) and lowest in RL (2.95). However, for methanotrophs, the average OTU number was the highest in CM (212–258) and lowest in the MF (134–265) and RL sites (170–214). The richness, as suggested by the ACE and Chao1, and the diversity, as suggested by the Shannon index, were highest in the CM site (Table S1).

3.3. Community Composition of Methanogens and Methanotrophs

3.3.1. Methanogens

Methanosarcinaceae, Methanoaetaceae, Methanobacteriaceae, Methanoregulaceae, Methanomassiliicoccaceae, Methanocellaceae, and Methanomicrobiaceae were detected in the soil samples (Fig.S2A), and the family Methanosarcinaceae was the most abundant of the methanogens in the three sampling sites (Figure 3A). However, the composition of methanogens varied significantly in different vegetation types. The relative percentage of Methanosarcinaceae was the highest in the RL site (75.25%), followed by the CM site (66.19%), and was the lowest in the MF site (41.32%). On the contrary, the relative percentage of Methanomassiliicoccaceae and Methanomicrobiaceae was the highest in the MF site (22.07% vs. 12.01%) and lowest in the RL site (0.74% vs. 0.22%). The family Methanobacteriaceae showed a consistent relative percentage among the three sampling sites (14.9–19.9%, Figure 3A).

The relative percentage of Methanosarcinaceae (56.09%, 37.56%, and 30.31% at depths of 0–10 cm, 10–20 cm, and 20–30 cm, respectively) and Methanobacteriaceae (29.33%, 8.30%, and 7.20% at depths of 0–10 cm, 10–20 cm, and 20–30 cm, respectively) were decreased with increasing depth in MF. However, the relative percentage of Methanomassiliicoccaceae (2.88%, 27.31%, and 36.03% at depths of 0–10 cm, 10–20 cm, and 20–30 cm, respectively) and Methanomicrobiaceae (2.73%, 16.42%, and 16.88% at depths of 0–10 cm, 10–20 cm, and 20–30 cm, respectively) were increased with increasing depth in MF. In the CM and RL sites, Methanosarcinaceae showed a relatively higher abundance in the 10–20-cm soil layer (80.67% and 80.39% in CM and RL, respectively) than in the 0–10-cm (53.81% and 73.06%, respectively) and 20–30-cm soil layers (64.10% and 72.30%, respectively, Figure 4A). In the CM site,
*Methanobacteriaceae* showed a relatively higher abundance in the 0–10-cm soil layer (40.08%) than that in the 10–30-cm soil layers (8.61–11.19%), but it showed relatively consistent abundance in the three soil layers in the RL site (14.46–19.89%, Figure 4A).

**Figure 3.** The relative percentage of Methanogens (A) and Methanotrophs (B) phylotypes at three sites in the Dongting floodplain. The data is average value across a 0–30 cm soil profile. Site abbreviations: MF: mudflat, CM: Carex meadow, RL: reed land.
3.3.2. Methanotrophs

The methanotroph groups detected in the three sampling sites included the Methylobacter, Methylosarcina, Methylogagea, Methylococcus, Methylocaldum, Methylocystis, Methylococcus, and Methylomonas genera (Figure S2B). The most common and dominant genus of methanotrophs in CM was Methylocystis (60.85%, type II methanotrophs, from the Rhizobiales order), the abundance of which was significantly higher than that in the MF (35%) and RL sites (18%). Methylobacter and Methylosarcina (type I methanotrophs, both from the Methylococcales order) were dominant in MF (51.00%) and RL (50.24%), respectively (Figure 3B).

The vertical patterns of the methanotroph community structure are shown in Figure 4B. The relative percentage of pmoA phylotypes (%) in MF was higher in the 10–30-cm soil layer (54.52–65.29%) than in the 0–10-cm layer (79.08%). However, the relative percentage of methanogen in RL was relatively constant among the three soil layers (47.23–53.22%, Figure 4B). The relative percentage of Methylocystis in CM was the highest in the 10–20-cm soil layer (33.18%). The relative percentage of Methylosarcina in RL was relatively constant among the three soil layers (47.23–53.22%, Figure 4B).
soil layer (79.08%). However, the relative percentage of *Methylosarcina* in RL was relatively constant among the three soil layers (47.23–53.22%, Figure 4B).

### 3.4. The Relationship between Methanogen and Methanotroph Communities and Environmental Factors

The soil environmental factors can be divided into two axes; the axes of RDA 1 (water regime condition, 80.31%) and RDA 2 (soil C and N availability, 14.65%) successfully explained the total variance for the methanogen–environment relationship, and the axes of RDA 1 (water regime condition) and RDA 2 (soil C and N availability) explained 81.29% and 13.87% of the total variance for the methanotroph–environment relationship, respectively (Figure 5A, B). WTD and pH were positively correlated with the *Methanosarcinaceae* distribution but negatively correlated with the distribution of *Methanomassiliicoccaceae*, *Methanosetaeaceae*, and *Methanomicrobiaceae*. MBC, MBN, TC, SOC, and TN were positively correlated with the *Methanobacteriaceae* distribution (Figure 5A, Table 2). For methanotroph communities, WTD and pH were positively correlated with the *Methylosarcina* but negatively correlated with the distribution of *Methylobacter* and *Methylocystis*. MBC, MBN, TC, SOC, and TN were positively correlated with the distribution of *Methylosinus* and *Methylococcus* (Figure 5B, Table 2). Since the SWC was significantly negatively correlated with the WTD and pH, the effects of SWC on methanogen and methanotroph communities were contrary to that of WTD and pH.

![Figure 5](image-url)

**Figure 5.** RDA ordination diagrams of methanogen (A) and methanotroph (B) community composition and environmental variables. MF, CM, and RL represent the three sampling sites of mudflat, *Carex* meadow, and reed land, respectively. Arrows point to variables associated with community composition, and the length of the arrows indicates the percentage of data explained by that variable. The following abbreviations indicate environmental factors: SWC: soil water content, MBC: microbial biomass carbon, MBN: microbial biomass nitrogen, SOC: soil organic carbon, TC: total carbon, TN: total nitrogen, WTD: water table depth.
β dance of methanotrophs indirectly through SOC and TN (total effect: 43.4%, indirect effect: 75.7%, total effect: 52.0%). SWC positively affected the abundance of methanotrophs (total effect: 17%).

Table 2. The coefficients of correlation (β) and determination (r²) between methanogen and methanotroph communities and environmental factors by redundancy analysis (Koh, #46).

| Environmental Factors | Methanogen RDA1(β₁) | Methanogen RDA2(β₂) | r² | Methanotroph RDA1(β₁) | Methanotroph RDA2(β₂) | r² |
|------------------------|----------------------|---------------------|----|----------------------|----------------------|----|
| WTD                    | 0.96                 | 0.29                | 0.54 *** | 0.99                 | 0.02                 | 0.94 *** |
| SWC                    | -0.86                | -0.52               | 0.35 ** | -0.99                | -0.02                | 0.26 * |
| pH                     | 0.96                 | 0.27                | 0.57 *** | 0.99                 | -0.03                | 0.80 *** |
| MBC                    | 0.69                 | -0.72               | 0.48 *** | 0.72                 | -0.69                | 0.21 * |
| MBN                    | 0.69                 | -0.72               | 0.36 ** | 0.30                 | -0.95                | 0.32 ** |
| SOC                    | 0.41                 | -0.91               | 0.40 ** | 0.22                 | -0.97                | 0.26 * |
| TC                     | 0.53                 | -0.84               | 0.35 ** | 0.33                 | -0.94                | 0.27 * |
| TN                     | 0.42                 | -0.90               | 0.34 ** | 0.31                 | -0.95                | 0.22 * |

WTD: water table depth; SWC: soil water content; MBC: microbial biomass carbon; MBN: microbial biomass nitrogen; SOC: soil organic carbon; TC: total carbon; TN: total nitrogen. Symbols *, **, and *** indicate at the significance levels of 0.05, 0.01, and 0.001, respectively.

3.5. Effect Pathways of Environmental Factors on CH₄-Cycling Microorganisms

The effect pathways of WTD, SWC, and pH on the abundance of methanogens and methanotrophs through soil C (SOC), soil N (TN), MBC, and MBN were successfully fitted by the piecewise SEM (Table 3). For methanogens, the WTD, SWC, and pH affected the abundance of mcrA directly and indirectly through MBC and MBN or the SOC, TN, MBC, and MBN (Figure 6A). Overall, the WTD and SWC positively affected the abundance of methanogens (WTD: direct effect: −28.0%, indirect effect: 43.4%, total effect: 15.4%; SWC: direct effect: −52.0%, indirect effect: 75.7%, total effect: 23.7%). However, pH negatively affected the abundance of methanogens (direct effect: 48.0%, indirect effect: −67.0%, total effect: −19%).

For methanotrophs, WTD positively affected the abundance of methanotrophs indirectly through the SOC and TN or the SWC (total effect: 34.8%). pH negatively affected the abundance of methanotrophs indirectly through SOC and TN (total effect: −51.0%). SWC positively affected the abundance of methanotrophs indirectly through SOC and TN (β = 0.64 × 0.99 × 1.01 = 0.64) but negatively affected the abundance of methanotrophs directly (β = −0.47, Figure 6B). Overall, SWC positively affected the abundance of methanotrophs (total effect: 17%).

Figure 6. Cont.
The abundance of methanogens was the highest in the CM sites, which may induce the highest CH$_4$ emission in the CM site at an intermediate elevation. However, the abundance of methanotrophs was significantly correlated with CH$_4$ emission in wetland soils. However, this conclusion still needs more evidence from other natural wetlands. In addition, the plant zonation sites can change from a CH$_4$ source to CH$_4$ sink with an increase in elevation in the Dongting Lake floodplain. The abundance of methanogens was the highest in the CM sites, which may induce the highest CH$_4$ emission in the CM site at an intermediate elevation. However, the abundance of methanotrophs was the highest in CM, indicating that CH$_4$ consumption may also be significant in CM sites. Fundamentally, soil CH$_4$ flux is responsible for the balance of CH$_4$ production and consumption by methanogens and methanotrophs, respectively [31]. We found that the ratio of methanogen/methanotroph was the highest in CM and lowest in RL, which can explain the significant CH$_4$ emission in CM and a noticeable CH$_4$ absorption in the RL sites (Figure 2). Chen et al. (2017) also reported a linear relationship between CH$_4$ emission and the methanogen/methanotroph ratio in the Chongxi wetland, a coastal restored wetland in eastern China [17]. Recently, Rey-Sanchez et al. (2019) also reported that the ratio of methanogens to methanotrophs in the upper portion of the peat was significantly correlated with CH$_4$ exchange velocity from a peat bog located in northeastern Ohio [32]. Therefore, we can infer that the methanogen/methanotroph ratio might be a critical factor in determining CH$_4$ emission in wetland soils. However, this conclusion still needs more evidence from other natural wetlands. In addition, the primer mb661r used in this study does not detect all the high-affinity methanotrophs [33], which might further explain the CH$_4$ sink function of the RL site.

**Figure 6.** Structural equation model (SEM) of the relationship between the abundance (qPCR) of methanogens (A) and methanotrophs (B) and environmental factors in the three sites. Solid red arrows represent positive paths ($p < 0.05$, piecewise s.e.m.), solid blue arrows represent negative path ($p < 0.05$, piecewise s.e.m.) and dotted blue arrows represent non-significant paths ($p > 0.05$, piecewise s.e.m.).

**Table 3.** Statistics of the structural equation model (SEM) of the relationship between the abundance of methanogens and methanotrophs and environmental factors in the three sites.

| Dependent Variables | Fisher.c | $p$ Value | AIC   | $R^2$ |
|---------------------|----------|-----------|-------|-------|
| methanogens         | 13.81    | 0.46      | 77.81 | 0.77  |
| methanotrophs       | 9.95     | 0.62      | 81.95 | 0.75  |

**4. Discussion**

**4.1. CH$_4$ Flux and Abundance of CH$_4$-Aycling Microorganisms**

In this study, both the CM and MF sites showed a significant CH$_4$ emission; however, the RL site exhibited a significant CH$_4$ absorption. This result indicated that the plant zonation sites can change from a CH$_4$ source to CH$_4$ sink with an increase in elevation in the Dongting Lake floodplain. The abundance of methanogens was the highest in the CM sites, which may induce the highest CH$_4$ emission in the CM site at an intermediate elevation. However, the abundance of methanotrophs was the highest in CM, indicating that CH$_4$ consumption may also be significant in CM sites. Fundamentally, soil CH$_4$ flux is responsible for the balance of CH$_4$ production and consumption by methanogens and methanotrophs, respectively [31]. We found that the ratio of methanogen/methanotroph was the highest in CM and lowest in RL, which can explain the significant CH$_4$ emission in CM and a noticeable CH$_4$ absorption in the RL sites (Figure 2). Chen et al. (2017) also reported a linear relationship between CH$_4$ emission and the methanogen/methanotroph ratio in the Chongxi wetland, a coastal restored wetland in eastern China [17]. Recently, Rey-Sanchez et al. (2019) also reported that the ratio of methanogens to methanotrophs in the upper portion of the peat was significantly correlated with CH$_4$ exchange velocity from a peat bog located in northeastern Ohio [32]. Therefore, we can infer that the methanogen/methanotroph ratio might be a critical factor in determining CH$_4$ emission in wetland soils. However, this conclusion still needs more evidence from other natural wetlands. In addition, the primer mb661r used in this study does not detect all the high-affinity methanotrophs [33], which might further explain the CH$_4$ sink function of the RL site.
4.2. CH₄-Cycling Microorganism Composition

4.2.1. Methanogen Composition in Three Vegetation Types

In this study, the Methanosarcinaceae was the most abundant family in the three vegetation types (Figures 3 and 4), indicating that the Methanosarcinaceae are the main CH₄-producing methanogens in the Dongting Lake floodplain. The relative percentage of Methanosarcinaceae was significantly increased in CM and RL sites, but the relative percentages of Methanomassiliicoccaceae and Methanomicrobiaceae (nonacetoclastic families) were decreased. This result indicated that the acetate pathway might be the dominant pathway for CH₄ production in CM and RL sites since plenty of acetate could be accumulated in the wetland soil due to root secretion and root anaerobic respiration along with plant development [15,34,35]. Furthermore, some nutrient materials (SOC, TN, NH₄-N, NO₃-N, etc.) produced by root decomposition can also serve as a substrate for Methanosarcinaceae methanogens, leading to a higher relative percentage of Methanosarcinaceae in the CM and RL sites, especially in the 10–20-cm soil layers (Figure 4A). These phenomena were also found in other freshwater wetlands, such as the freshwater–brackish water gradients of the Min River [24], Zoige peatland, and Sanjiang Carex lasiocarpa marsh [36]. However, besides the Methanosarcinaceae, the Methanomicrobiaceae and Methanobacteriaceae are prevalent in MF, suggesting that an H₂/CO₂ pathway might be the main process for CH₄ production in the frequently flooded MF site. These results were also confirmed by Kemnitz et al. (2004), who reported that Methanomicrobiaceae was only found in the permanently and frequently flooded soils in a riparian flooding gradient in the Netherlands [14].

Interestingly, the family Methanomassiliicoccaceae, which was recently discovered in human and animal gastrointestinal tracts (GITs) [37], was detected in the MF and CM sites in our study. One possible explanation for this is that Dongting Lake is an important wintering ground for migratory birds, and CM and MF sites are the main places for their feeding and inhabitation [38]. In this context, the inhabitation of migratory birds in the Dongting Lake floodplain might provide a positive feedback for CH₄ production.

4.2.2. Methanotroph Composition in Three Vegetation Types Along an Elevation Gradient

The dominant methanotrophic genera detected in our study were Methylobacter, Methylosarcina (both belonging to type I methanotrophs), and Methylocystis (belonging to type II methanotrophs, Figure 3B). We found that Methylocystis represented the dominant methanotrophic bacteria in the CM site. However, Methylobacter and Methylosarcina were the dominant genera in the MF and RL sites, respectively. It has been documented that type I and type II methanotrophs generally have distinct life strategies, enabling them to predominate under different conditions and maintain functionality [39,40]. Generally, Type I methanotrophs, which respond rapidly to substrate availability and exist as the predominantly active community in many environments, can thus be classified as competitors (C) and competitors-ruderals (C-R). On the contrary, type II methanotrophs were found to persist in inactive states and show versatility in substrate utilization; they are classified as stress tolerators (S) and stress tolerator-ruderals (S-R) [40]. Therefore, the genus Methylocystis (type II methanotrophs), which has repeatedly been found to be associated with a wide variety of environments (such as rice paddies, different upland and hydromorphic soils, landfills, peatlands, and glacier fore fields) [41], was not only dominant in the CM site but also prevalent in the MF and RL sites in the Dongting Lake floodplain (Figure 3B). However, because of the frequent flooding disturbance and water shortage in MF and RL, the methanotrophs living in MF and RL should be more competitive for acquiring O₂ and/or water. Therefore, Methylobacter and Methylosarcina (both are type I methanotrophs) act as competitors in MF and RL sites, respectively.

Further, the SWC may act as a critical factor for methanotroph distribution in different soil layers, since the SWC determines soil aeration along a soil profile [26]. In the MF site, the SWC was significantly higher in the 10–30 cm soil layer than in the 0–10 cm soil layer. In the CM site, the SWC was higher in the 0–10 cm and 20–30 cm soil layers, while in the RL site, the SWC was relatively consistent among
the three soil layers (Table S2). According to the C-S-R functional classification of methanotrophs, the relative percentage of the genus *Methylobacter* in MF should be increasing for O\textsubscript{2} competition with an increase in soil depth, while the relative percentage of the genus *Methylosarcina* was consistent in the RL site. Moreover, the diversity of nitrogen metabolism in methanotrophs is well documented [41–43]. Type II methanotrophs and members of the type I genus *Methyllococcus* have been shown to be capable of nitrogen fixation, while other type I methanotrophs are not capable of performing this process [44]. Therefore, other factors, such as soil ammonium and nitrate, may also play important roles in selecting the methanotroph type in nature wetland soils, which should be considered in future studies.

**4.3. Environmental Controls on CH\textsubscript{4}-Cycling Microorganisms**

Redundancy analysis used multiple linear regressions to explain the variation between independent and dependent variables, and these calculations are performed within the iterative procedure to find the best ordination of the objects [45]. The results of the RDA analysis indicated that the WTD, SWC, and pH are critical factors influencing the methanogen and methanotroph community structure. From water-adjacent areas to higher-elevation lands, the WTD increased, but the SWC decreased, and the soil environment may change from anaerobic to aerobic. Further, the soil pH may also increase with a decrease in SWC [46]. These events may change the living conditions for CH\textsubscript{4}-cycling microorganisms and eventually reduce CH\textsubscript{4} emission with an increase in elevation in a floodplain wetland [16,47].

Further, SEM analysis also showed that the WTD and SWC both had negative direct effects but positive indirect effects on the abundance of methanogens through the soil C and N concentration (i.e., SOC and TN), MBC, and MBN. In the Dongting Lake floodplain, the Carex and reed communities were distributed at 24–25 m and 25–28 m elevations, respectively, and the WTD was 1–1.5 m and 2–3 m in the Carex and reed communities, respectively (Table S2). On the one hand, with an increase in elevation, because of the plant growth and litter decomposition, the soil C and N concentration and microbial biomass were significantly increased, which can supply sufficient substrate for methanogen growth [7,16]. On the other hand, the continuous increase in elevation induced SWC decline; more oxygen can permeate into the soil profile through oxygen loss from rhizosphere [47], which may inhibit methanogen growth in the RL site. Moreover, due to a slower decomposition rate in RL, the complex organic compounds in the RL site cannot be directly utilized by methanogens, which may also result in a lower abundance of methanogens in the RL site [48,49]. Similarly, the continuous decrease in elevation induced SWC increase, but soil C and N shortage may cause a lower methanogen abundance in the MF site. Generally, methanogens are usually active over a pH range of 6.6–7.8, the optimum pH being 6.8, whereas most methanogens can be inhibited if the pH is lower than 6.1 or higher than 8.3 [50,51]; however, methanogen activity was occasionally found to be very high in considerably more acidic environments in ombrotrophic peats [6]. The soil pH in our study site ranged from 7.51–8.07 (Table S2). Therefore, the soil pH could have negatively affected methanogen abundance through its negative influence on SOC and MBC, as confirmed by our SEM results.

For methanotrophs, a higher SOC and TN content can facilitate methanotrophs to assimilate gaseous methane in the soil profiles [52]. The WTD and SWC also had positive indirect effects on the abundance of methanotrophs through their positive effects on the SOC and TN, which can explain the significantly higher abundance of methanotrophs in the CM site, especially in 0-10-cm soil layers. Additionally, more oxygen in the surface layers, combined with moderate SWC, also favors methanotroph growth (Table S2). Similar to the effect of the soil C and N on methanogens, the complex organic compounds in the RL site may also inhibit methanotroph growth, although there is an aerobic environment in the RL site. The soil pH had negative indirect effects on the abundance of methanotrophs, which suggested that methanotrophs may favor a neutral and/or relatively acidic environment [53]. Factually, different methanotrophs are known to have different pH optima [54], and pH can only be considered an indicator rather than a decisive factor for methanotroph abundance in this study.
Previous studies have shown emissions of CH\(_4\) from northern peatlands vary as a function of temperature, pH, substrate and nutrient availability, etc.\[4\]. However, in the subtropical floodplains, the emissions of CH\(_4\) combined with methanogens/methanotrophs composition are mainly influenced by WTD, SWC, and pH. The significant change in WTD, SWC, and pH not only determines soil redox potential for CH\(_4\) production but also controls substrate and nutrient availability by influencing biogeochemical cycles. Besides the environmental factors considered in this study, other environmental variables, such as the oxygen availability and concentration of metal ions (e.g., Fe\(^{3+}\), Mn\(^{4+}\), and SO\(_4^{2-}\)) in the soil profile and different vegetation, are also reported as critical factors influencing CH\(_4\) emission and methane-cycling microorganisms [55–58]. These observations may account for the unexplained variance of methanogen/methanotroph–environment relationship in this study. This needs further study in the future.

5. Conclusions

This study provided fundamental insights into soil CH\(_4\) emissions, the community structure of CH\(_4\)-cycling microorganisms, and their environmental controls in a subtropical inland floodplain. The Dongting Lake floodplain may change from a CH\(_4\) source to a CH\(_4\) sink with vegetation succession along an elevation gradient, and the methanogen/methanotroph abundance ratio can be used as a CH\(_4\) emission proxy in natural wetlands. In the Dongting Lake floodplain, Methanosarcinaceae is the main CH\(_4\)-producing methanogen, Methylocystis (type II methanotrophs) represents the dominant methanotrophic bacteria in CM, and Methylobacter and Methylosarcina (type I methanotrophs) are the dominant genera in MF and RL. Methanogen and methanotroph community structures are mainly controlled by the WTD, SWC, and pH along the elevation gradient. The continuous increase in reed area combined with the decrease in CM, which is caused by the decrease in WTD, may mitigate CH\(_4\) emission and enhance the CH\(_4\) sink function during the non-flood season in the Dongting Lake floodplain.

Supplementary Materials: The following are available online at http://www.mdpi.com/2073-4433/11/9/997/s1, Figure S1: Rarefaction curve for mcrA(A) and pmoA(B) sequences clustered at 97% similarity across the three site and soil depths in the Dongting floodplain. Figure S2: The phylogenetic trees of methanogens (A) and methanotrophs (B) using approximately-maximum-likelihood method on the basis of the mcrA and pmoA sequences in the soils of three sites in the Dongting floodplain, Table S1: Richness and diversity indices of methanogen (mcrA) and methanotroph (pmoA) at different soil depths in the Dongting Lake floodplain. Table S2: Physicochemical properties of soils at each sampling site on November 10, 2016, in the Dongting Lake floodplain.

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