1. INTRODUCTION

Aquaculture is one of the most rapidly growing traditional agricultural industries, providing approximately half of the fish consumed by humans (Lu et al. 2020). The requirement for aquatic products has supported the fast expansion of aquaculture, which has also led to huge economic benefits (Zhao et al. 2020). Among aquacultured resources, Urechis unicinctus is one of the most valuable products in Asia due to its high nutritional and medicinal values. U. unicinctus has the capacity to tolerate and utilize sulfide, a toxicant (Shi et al. 2012); this capacity has ecological value for improving the biological suitability of sulfur-containing substrates in coastal substrates. Polyculture systems can improve biological diversity, enhance the utilization of input material, strengthen the self-purifying ability of aquaculture water, improve economic efficiency, and reduce environmental pollution. U. unicinctus are always polycultured with Penaeus japonicus or Stichopus japonicus (Yuan et al. 2015), and can make full use of a large amount of
Intestinal microorganisms are an important resource pool for marine organisms and previous studies have mainly focused on fish (Mouchet et al. 2012, Zheng et al. 2018, Parris et al. 2020). Therefore, intestinal microorganisms and their interactions with the host have become research ‘hotspots’ in the fields of microbiology and medicine.

Intestinal microbes have some unique metabolic pathways, and they can establish complex metabolic networks with the host; it is therefore of great significance to clarify the microbial diversity and functions in the intestinal environment of U. unicinctus. For this study, we selected U. unicinctus from 2 different habitats: (1) in pond polyculture with P. japonicus and (2) a coastal intertidal flat to investigate intestinal microbial diversity and conduct a functional analysis by 16S rDNA 454 high-throughput sequencing technique. The goal of this research was to (1) assess the difference in bacterial communities between the 2 different U. unicinctus habitats and screen out discriminatory taxa, (2) analyze intestinal microbial function, and (3) evaluate the correlation between gut microbiota and the bacterial communities in the 2 habitats.

2. MATERIALS AND METHODS

2.1. Animals and sampling

Urechis unicinctus with a mean fresh mass of 29.0 ± 4.4(SD) g were collected from a coastal intertidal flat around Furong Island, Laizhou Bay, Yantai, China. The worms occupied burrows below the low-tide line, at a depth of 4 m. Living worms were maintained in containers with clean seawater for approximately 2 h before being transported to the laboratory. Upon arrival at the lab, the worms were euthanized and dissected under sterile conditions, and the intestinal contents were collected in sterilized storage tubes and stored at −80°C for later use. Water and sediment samples were collected from the surrounding environment, frozen immediately in a liquid nitrogen tank, and then stored at −80°C for later use. The worms, water, and sediment samples from the coastal flat are labeled SUU, SW, and SS, respectively.

U. unicinctus with a mean fresh mass of 16.6 ± 5.0 (SD) g were collected from a seawater aquaculture pond polycultured with Penaeus japonicus, at a depth of 1.0–1.5 m. The pretreatment procedures were the same as described above. The worms, water, and sediment samples from the pond polyculture are labeled PUU, PW, and PS, respectively.

2.2. DNA extraction and 16S rDNA Amplicon library construction

DNA was extracted from the intestinal contents using a FastDNA SPIN Kit for Soil (MP Biomedicals, Santa Ana, CA). A total of 150 ng of DNA was amplified using a 16S rDNA V3–V4 region primer set (5′-TCGTCGGCAGCGTCAGATGTGTATAAGAGACAGCAGCAGCCGGTCTTAAACCCAGTCAC-3′ and 5′-CAAGCAGAAGACGGCATACGAGAT-3′) for U. unicinctus and a 16S rDNA V4 region primer set (5′-CCTACGGGNGGCWGCAG-3′ and 5′-GACTACHVGGGTATCTAATCC-3′) for P. japonicus. The PCR products were purified using a QIAquick PCR Purification Kit (Qiagen, Venlo, Netherlands) and quantified using a Quant-iT PicoGreen dsDNA Assay Kit (Thermo Fisher Scientific, Waltham, MA). The PCR products were poolsequencing technique. The goal of this research was to (1) assess the difference in bacterial communities between the 2 different U. unicinctus habitats and screen out discriminatory taxa, (2) analyze intestinal microbial function, and (3) evaluate the correlation between gut microbiota and the bacterial communities in the 2 habitats.

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2.2. DNA extraction and PCR amplification

We used an E.Z.N.A.® soil DNA Kit (Omega Bio-Tek) to extract the total DNA of all the samples, following the manufacturer's protocols. A NanoDrop 2000 UV-vis spectrophotometer (Thermo Scientific) was employed to determine the concentration and purification of final DNA, and 1% agarose gel electrophoresis was applied to check the DNA quality. Using a thermocycler PCR system (GeneAmp 9700), the primers 338F and 806R (Chen et al. 2020) were used to amplify the V3–V4 hypervariable regions of the 16S rRNA gene of all samples. The PCR reactions were conducted according to the literature (Zhao et al. 2019, Xu et al. 2020), and the PCR products were excised from a 2% agarose gel. An AxyPrep DNA Gel Extraction Kit (Axygen Biosciences) was used for further purification, and DNA was quantified using Quanti-Fluor™-ST (Promega).

2.3. Sequencing on the Illumina MiSeq platform

The purified amplicons were submitted to the Majorbio Bio-Pharm Technology Co. Ltd. (Shanghai, China) following standard protocols. They were then pooled in equimolar amounts and paired-end sequenced (2 × 300) on an Illumina MiSeq platform (Zhao et al. 2019, Xu et al. 2020).

2.4. Bioinformatic data analysis

Raw fastq files were quality-filtered by Trimmomatic and merged by FLASH using UPARSE (Zhao et al. 2019, Xu et al. 2020, Kong et al. 2021). Based on the Silva (SSU123) 16S rRNA database, each 16S rRNA gene sequence was analyzed by the RDP Classifier algorithm (http://rdp.cme.msu.edu/), with a confidence threshold of 70%. Richness and diversity, the community structure, and functional analysis were performed using the free online Majorbio Cloud Platform (http://www.majorbio.com).

KEGG orthology (KO), cluster of orthologous groups (COG), and pathways of the samples from amplicon sequencing results were predicted using the Phylogenetic Investigation of Communities by Reconstruction of Unobserved States (PICRUSt) based on the Greengenes 13.5 database (Langille et al. 2013). All software packages were freely available on the Majorbio Cloud Platform.

2.5. Data analysis

The relationship between the selected taxonomical groups (abundant phyla, genera, classes, orders, or families), the observed operational taxonomic units (OTUs), and the bacterial community index were calculated using SPSS 18.0 software. All statistical analyses were conducted using SPSS software (version 18.0). A value of $\alpha = 0.05$ was accepted as the criterion for statistical significance. The homogeneity of variance of all data was tested by Levene's test and the normality of residuals was tested by Shapiro-Wilk test. One-way ANOVA was employed to conduct multiple-group comparisons, and Student's $t$-test (2-tailed test) was used to conduct 2-group comparisons. Data are presented as means ± SD.

3. RESULTS

The raw reads were deposited into the NCBI Sequence Read Archive (SRA) database under accession number PRJNA681016. A total of 902 871 optimized sequences were obtained, from 2 intestinal samples, 2 water samples, and 2 sediment samples (1 sample each from the coastal zone and the polyculture pond, with 3 replicates for each sample) (Table 1). The numbers of clean sequences in the 2 different habitats and *Urechis unicinctus* gut were significantly different; the number of sequences in the pond samples was much lower than that from the coastal samples. After rarefaction to equal sequencing depth, a total of 4945 OTUs were obtained from all samples. The OTU numbers in SUU, PUU, SS, PS, SW, and PW were 513 ± 77, 618 ± 258, 1738 ± 68, 1783 ± 79, 1074 ± 168, and 445 ± 21, respectively. The OTU numbers in the sediment samples were larger than in the water samples and the gut of *U. unicinctus*. We identified 52 OTUs that were shared among all samples, and only a few unique OTUs (32) were found in PW samples, indicating that relatively few microorganisms inhabited PW. Intestine samples of *U. unicinctus* living in the 2 habitats had only 66 OTUs in common, suggesting significant differences in the gut bacteria between the 2 habitats (see Figs. S1 & S2 in the Supplement at www.int-res.com/articles/suppl/q013p211_supp.pdf).

3.1. Richness and diversity analysis

The richness and diversity of the microbial community can be studied through alpha diversity analysis,
including a series of indices used to estimate species abundance and diversity of the environmental community (Table 1). Abundance-based coverage estimators (ACE) and Chao indices indicate the community richness of the samples (see Text S1 in the Supplement for details of how these indices were calculated). In both habitats, species richness was highest in the sediment samples. The gut bacterial community richness of *U. unicinctus* living in the pond was higher than that of *U. unicinctus* inhabiting the mudflat. The bacterial community richness of SW was much higher than that of PW; the ACE and Chao values of the SW samples were 1969 ± 208 and 1655 ± 143, nearly 2.5 times those of the PW samples. Shannon and Simpson indices reflect community diversity. Except for the PW, the Shannon and Simpson indices showed no significant differences between the gut and habitat samples. Good’s coverage index of all samples was >0.98, indicating that the OTUs of each sample were well captured.

### 3.2. Bacterial composition of the intestine and the habitat

We identified 58 different bacterial phyla from all samples. The dominant bacterial phyla that appeared in all samples were *Proteobacteria, Bacteroidetes, Firmicutes, Actinobacteria, Cyanobacteria, Chloroflexi*, and *Acidobacteria* (Figs. 1a & S3). Among all samples, the abundance of *Proteobacteria* in SUU was the highest, and the abundance of *Bacteroidetes* was

| Sample ID | Effective sequences | OTUs | ACE | Chao | Shannon | Simpson | Good’s coverage |
|-----------|---------------------|------|-----|------|---------|---------|----------------|
| SUU       | 56063 ± 2473        | 513 ± 77 | 521 ± 75 | 526 ± 70 | 4.43 ± 0.22 | 0.052 ± 0.013 | 0.9997 |
| PUU       | 52192 ± 4670        | 618 ± 258 | 628 ± 259 | 643 ± 261 | 4.85 ± 0.27 | 0.036 ± 0.006 | 0.9995 |
| SS        | 47977 ± 3739        | 1738 ± 68 | 2190 ± 56 | 2138 ± 57 | 4.46 ± 0.06 | 0.087 ± 0.006 | 0.983 ± 0.006 |
| PS        | 44515 ± 2865        | 1783 ± 79 | 2225 ± 83 | 2204 ± 60 | 4.42 ± 0.15 | 0.085 ± 0.014 | 0.986 ± 0.001 |
| SW        | 52367 ± 3622        | 1074 ± 168 | 1969 ± 208 | 1655 ± 143 | 4.44 ± 0.10 | 0.039 ± 0.016 | 0.990 ± 0.003 |
| PW        | 47943 ± 2161        | 445 ± 21 | 877 ± 106 | 676 ± 54 | 3.44 ± 0.02 | 0.066 ± 0.001 | 0.995 |

Table 1. Richness, diversity, and coverage estimates obtained at genetic distances of 3%. Data are presented as means ± SD. SUU (PUU): gut samples of *Urechis unicinctus* collected from the coast (collected from a pond polyculture); SS (PS): sediment samples collected from the coast (pond polyculture); SW (PW): water samples collected from the coast (pond polyculture); OTUs: operational taxonomic units; ACE: abundance-based coverage estimators.
Tang et al.: Intestinal microbial diversity of Urechis unicinctus.

Firmicutes were not found in the water samples, but were abundant in the sediment samples, whereas Actinobacteria were much higher in the water samples than in the other samples. Cyanobacteria were found almost exclusively in the SW samples. Chloroflexi and Acidobacteria were present in sediment samples (Fig. 1b).

Bacterial community composition in the intestine and the habitat samples at the genus level is displayed in Fig. 2, with a total of 1295 genera identified in all samples. The genus with the highest relative abundance in SUU (26.78%) and PUU (22.10%) was Acinetobacter. norank_f_Cryomorphaceae\(^1\) accounted for 6.70% of all genera in PUU, while this relative abundance declined sharply in SUU (0.14%). Psychrobacter and Planococcus were the major genera in SS and PS, accounting for 41.48 and 42.30% of all genera. The relative abundance of genera in the water samples presented a significant difference from the other samples. norank_f_norank_o_Chloroplast (14.48%), Lentibacter (9.24%), Pseudoalteromonas (7.07%), and Candidatus_Aquiluna (5.39%) were the top 4 genera in SW. Smaller percentages of Planktomarina (17.04%), Candidatus_Aquiluna (12.65%), Vibrio (9.28%), Pseudoalteromonas (9.24%), and norank_f_Microbacteriaceae (7.44%), NS3a_marine_group (4.74%), unclassified_f_Rhodobacteraceae (4.42%), and norank_f_Cryomorphaceae (4.10%) were found in PW.

Additionally, the phylogenetic relationships of these genera differed (Fig. 2). Most genera belonged to Proteobacteria, including Psychrobacter, Candidatus_
Puniceispirillum, Pseudoalteromonas, Lentibacter, Acinetobacter, and Pseudomonas. Actinobacteria includes norank_f_norank_o_Actinomarinales, Candidatus_Actinomarina, Candidatus_Aquiluna and norank_f_Microbacteriaceae. Planococcus, Exiguobacterium and Staphylococcus belong to Firmicutes. norank_f_Cryomorphaceae and NS3a_marine_group are members of Bacteroidetes.

The relationship between samples and species on the genus level is presented in a Circos diagram (Fig. 3). Acinetobacter was mainly found in the intestinal samples, and Psychrobacter and Planococcus
were key species in the sediment samples. The genera found in water samples were diverse. The results were consistent with those shown in Fig. 2.

3.3. Bacterial community analysis and comparison

To compare bacterial community composition of the gut microbiota of *U. unicinctus* to those of their environments, we used principal co-ordinates analysis (PCoA) of weighted UniFrac distances (Fig. 4a). The 18 samples were separated into 3 clusters according to the second principal component (PC1) axis, accounting for 41.94% of the total variance. The gut microbiota (SUU and PUU) belonged to one group and were significantly different from the sediment samples (SS and PS) and the water samples (SW and PW). These groups are also consistent with the sample collection areas. Based on the UPGMA clustering tree (Fig. 4b), the taxonomic composition of all samples revealed 3 different groups, which was consistent with the results of the PCoA analysis. The gut microbiota (SUU, PUU) were clustered into one set in Fig. 4b. ANOSIM of all samples at the genus level also reflected that the species in intestinal samples were quite different from those in water and sediment samples (Fig. S4).

To detect species that exhibited differences in abundance in different microbial communities, the 15 top genera were selected to assess the significance of the observed differences in the intestine of *U. unicinctus* and the 2 different habitats using 1-way ANOVA (Fig. 5a). The abundances of *Psychrobacter* and *Planococcus* were extremely high in the sediment samples, which were significantly different from the other 2 groups. The genus *Acinetobacter* was only found in the intestine of *Urechis unicinctus*. *norank_f_Cryomorphaceae* was also found in PUU. The genera in water samples were significantly different from the other 2 groups. *Planktomarina*, *Candidatus_Aquiluna*, *norank_f_Cryomorphaceae*, and *Vibrio* were abundant in PW samples. The relative abundance of *norank_f_norank_o_Chloroplast* and *Lentibacter* was much higher in the SW samples. *Pseudoalteromonas* showed no obvious difference between SW and PW. To explore the different species inhabiting the intestines of *U. unicinctus* from the 2 different habitats, we again selected the 15 top genera to assess significance using Student’s *t*-test (Fig. 5b). Except *norank_f_Cryomorphaceae*, the
Fig. 5. Analysis of species differences at the genus level. Data are presented as means. (a) Species differences in samples from the intestine of *Urechis unicinctus* and the 2 different habitats (group abbreviations as in Table 1) using 1-way ANOVA (*0.01 < p ≤ 0.05; **0.001 < p ≤ 0.01; ***p ≤ 0.001). (b) Species differences in the intestine of *U. unicinctus* from 2 different habitats using Student's *t*-test (2-tailed test; *p ≤ 0.05).
other 14 top genera showed no significant differences between the 2 intestinal samples. PUU and SUU had more species in common than the other samples did, indicating that the relationship between the 2 intestinal samples was much closer (Fig. S5). The results indicated that *U. unicinctus* can survive and grow well regardless of whether it is maintained in pond polyculture or in coastal mudflats.

### 3.4. Functional analysis of intestinal microbial community

COG analysis classified all intestinal microbial genes into 23 functional categories (Fig. 6) (Table S1). In this classification, except for Group S (‘function unknown’), the largest group involved in the general function was amino acid transport and metabolism (Group E: 10.05% SUU, 10.08% PUU), which was followed by translation, ribosomal structure, and biogenesis (Group J: 6.76% SUU, 7.32% PUU); energy production and conversion (Group C: 6.97% SUU, 6.88% PUU); cell wall/membrane/envelope biogenesis (Group M: 6.37% SUU, 6.73% PUU); inorganic ion transport and metabolism (Group P: 6.81% SUU, 6.46% PUU); transcription (Group K: 5.99% SUU, 5.80% PUU); and carbohydrate transport and metabolism (Group G: 5.02% SUU, 5.15% PUU). Details are provided in Table S1.

The functional categories (KEGG level 2) of bacterial communities as predicted by PICRUSt analysis through a heatmap are shown in Fig. 7 & Table S2. The metabolic pathways were the largest pathways (68.56% PUU, 67.93% SUU). Following metabolism, the important functions of gut microbiota were genetic information processing (9.64% PUU, 8.85% SUU) and environmental information processing (8.36% PUU, 9.31% SUU). The highest relative abundance of genes were related to carbohydrate metabolism (Table S3), including pyruvate metabolism, glyoxylate and dicarboxylate metabolism, amino sugar and nucleotide sugar metabolism, and fructose and mannose metabolism (Table S4, Fig. S6). In addition to carbohydrate metabolism, the genes for amino acid metabolism accounted for 11.7% of the total genes, including the genes to metabolize arginine, proline, glycine, serine, threonine, phenylalanine, tyrosine, alanine, aspartate, glutamate, cysteine, methionine and histidine. Interestingly, except for nitrogen metabolism and oxidative phosphorylation, methane metabolism was one of the main functions of energy metabolism (Fig. S7). The gut intestinal microbiota could also metabolize nucleotides (purine and pyrimidine), cofactors and...
vitamins, lipid, xenobiotics, terpenoids, polyketides, and glycan. Translation, replication, and repair and folding, sorting, and degradation were the main functions related to genetic information processing, with a relative abundance of approximately 9.2%. The bacterial secretion system was of great significance for membrane transport, which was helpful for the gut microbiota to access to environmental information together with the signal transduction system, such as the ABC transporter. Quorum sensing is a critical function for cellular communication in prokaryotes. Bacterial chemotaxis plays an important role in cell motility. The 2 functions above associated with cell growth and death were the top 3 functions of cellular processes (6.4%) at KEGG level 2. The genes for cationic antimicrobial peptide (CAMP) and vancomycin resistance were also found to relate to antimicrobial drug resistance. Platinum drug resistance and antifolate, which represent antineoplastic drug resistance, were among the functions of gut microbiota in *U. unicinctus*.

4. DISCUSSION

In this work, we first investigated the intestinal microbial community of *Urechis unicinctus* living in 2 different habitats. Compared to the studies of sea cucumbers (Zheng et al. 2018), the number of total effective OTUs in this study was much larger. We
selected 18 samples, including 6 gut microbiota, to amplify the prokaryotic 16S rRNA gene, and 902,871 pyrosequencing reads in total were generated. We identified 445-1783 prokaryotic OTUs at a 93% sequence similarity level (Table 1).

Proteobacteria have a competitive edge in surviving in various ecological niches due to their variable morphology and versatile physiology. The relative abundance of Proteobacteria in the oceans was 57.9% (Shin et al. 2015), so they are a major phylum in intestinal samples of many marine organisms, including sea cucumbers (Zhang et al. 2018). Proteobacteria was the dominant phylum in the intestine of the sea cucumber Holothuria glaberrima (~50%) (Weigel 2020). In our research, Proteobacteria were dominant in all samples, including intestinal samples of U. unicinctus, seawater, and sediment from 2 habitats. Bacteroidetes was the phylum with the second highest relative abundance in all samples. Bacteroidetes are able to produce extracellular enzymes with degradative capabilities (Huang et al. 2020), and they might contribute to carbohydrate metabolism (Weigel 2020). In this study, the abundance of Bacteroidetes was stable in the intestinal samples (SUU and PUU), indicating that the intestinal bacteria have adapted to the environmental stresses and help the host to decompose and metabolize the diet (Duan et al. 2020). However, the relative abundances of Firmicutes were almost undetectable in 2 water groups. The abundance of Firmicutes was higher in sediments than in the gut (Fig. 1b). Firmicutes, which include a number of beneficial bacteria, could be more effective for extracting energy from the diet, and they can prevent the production of inflammatory cytokines and pathogen-induced intestinal function disruption in fish (Duan et al. 2020). Bacteroidetes participate in the degradation of different biological polymers, such as cellulose, chitin, and pectin (Williams et al. 2013). Previous studies have demonstrated that an elevated ratio of Firmicutes to Bacteroidetes is associated with weight gain and metabolic disorders (Zhao et al. 2019). Therefore, the stable Firmicutes/Bacteroidetes ratio of intestinal samples (SUU and PUU) indicates that energy metabolism of U. unicinctus might be healthy. Due to their ability to produce various natural drugs, enzymes, and bioactive metabolites (Jiang 2013), Actinobacteria are promising potential probiotics in aquaculture (Das et al. 2008). Cyanobacteria presented a high relative abundance only in the SW samples.

At the genus level, several dominant genera exhibited large differences among groups. Psychrobacter had the highest relative abundance in the SS samples. Psychrobacter is one of the dominant taxa in low-temperature environments that has the ability to degrade PAHs, is heavy-metal-resistant, and is presumably capable of denitrification in psychrophilic environments (Lasek et al. 2017). Acinetobacter represented over 20% of the relative abundance of the gut microbiota of SUU and PUU. Acinetobacter exhibit antibacterial activity against some tested human and fish pathogenic bacteria (Jami et al. 2015), and they are also resistant to many drugs (Wagenvoort et al. 2002), with defense mechanisms including efflux pumps, outer membrane porins, lipopolysaccharides, degradation enzymes (β-lactamases, metalloproteinases, penicillin-binding proteins), and biofilm formation (Bonomo & Szabo 2006, Livemore & Woodford 2006). Few Acinetobacter can cause infectious diseases (de Breij et al. 2010), although researchers have verified that this genus might infect marine fishes and increase the threat of transmission from fishes to humans (Anand & Suthindhiran 2020). The relative abundance of norank_f_Cryomorphaceae in PUU was significantly higher than that in SUU. Although little is known about this genus, its capacity to degrade proteins and algal storage polysaccharides has been confirmed (Grieb et al. 2020).

The physiological functions of hosts are greatly influenced by their intestinal microbiota (Goldsmith & Sartor 2014). Except the ‘function unknown’ group, the largest group involved in the general function was amino acid transport and metabolism, followed by translation, ribosomal structure and biogenesis, energy production and conversion, cell wall/membrane/envelope biogenesis, inorganic ion transport and metabolism, transcription and carbohydrate transport and metabolism based on the COG analysis. This is consistent with the literature (DeWitt & Kudsk 1999).

Based on the KEGG analysis, the largest group was amino acid and carbohydrate metabolism, indicating that intestinal microbes might prefer a diet rich in carbohydrates and proteins. Metabolism was the main function of the gut microbiota in U. unicinctus. In addition to nutrient metabolism, they also had capacity to degrade some harmful substances in the host including xenobiotics such as terpenoids and polyketides. Gut microorganisms can provide energy from digestion and metabolism based on multiple energy metabolism pathways (Webster 1980), such as nitrogen, oxidative phosphorylation, and methane metabolism.

Membrane transport and signal transduction are important for bacteria to access environmental information and conduct biological processes (Cornejo-Granados et al. 2017). The microbial genes involved
in membrane transport and signal transduction were abundant in the gut samples from *U. unicinctus*. These functional genes could also be found in tilapia (Wu et al. 2020). The genes encoding CAMP, vancomycin, platinum drug resistance, and antifolate resistance were also found among gut microbiota to tolerate the pathogenic bacteria and drugs, which indicated that the gut microbiota could modulate and enhance the immune system of *U. unicinctus* along with Th17 cell differentiation and the IL-17 signaling pathway (Belkaid & Hand 2014). Some other important pathways, including glucagon, insulin, estrogen, and adipocytokine signaling pathways and thyroid hormone synthesis and signaling pathways, suggested a vital role of the intestinal microbiota in the modulation of the endocrine system. The genes encoding GABAergic synapse (ko04727) and glutamatergic synapse (ko04724) were included in the KEGG analysis of gut microbes on level 3 (Table S4). Bacterial chemotaxis, flagellar assembly, quorum sensing, and biofilm formation could support the gut microorganism moving and communicating among the cellular processes to adapt to the intestinal environment of the host. These results may be of reference value to the poly-aquaculture management of *U. unicinctus* with other aquaculture organisms.

5. CONCLUSION

This study represents the first high-throughput absolute abundance quantification analysis of the gut microbiota diversity and functions in *Urechis unicinctus* from 2 different habitats. We found that *Proteobacteria, Bacteroidetes, Firmicutes, and Actinobacteria* were the dominant phyla in the gut microbial community, and the top genera were *Acinetobacter*, *Ralstonia*, and *Brevundimonas*. *Psychrobacter* and *Planococcus* were significantly high in sediment samples, while in sea water samples, *Planktomarina, norank_f_norank_o_Chloroplast*, and *Candidatus_Aquiluna* were significantly high in the gut microbiota. The gut microbiota were significantly different from the SS and SW samples. The major composition of gut bacteria did not show significant differences among the top 15 genera except *norank_f_Cryomorphaceae* in PW and SW.

The functions of intestinal microbial community largely centered on metabolism. In addition to the main carbohydrate and amino acid metabolism, energy metabolism including nitrogen metabolism, oxidative phosphorylation, and methane metabolism were relatively important. The gut microbiota could access information from the intestinal environment by virtue of membrane transport and signal transmission systems. The findings in the present study provide an understanding of the gut microbiome composition and diversity in *U. unicinctus* from 2 different habitats. We found no significant differences in the gut microbiota, indicating that *U. unicinctus* can survive and grow both in pond polyculture as well as in coastal mudflats, which can be instructive for the management of animal health in *U. unicinctus* farming activities, and for further gut microecology research.

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