Diversity, function and assembly of mangrove root-associated microbial communities at a continuous fine-scale

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Mangrove roots harbor a repertoire of microbial taxa that contribute to important ecological functions in mangrove ecosystems. However, the diversity, function, and assembly of mangrove root-associated microbial communities along a continuous fine-scale niche remain elusive. Here, we applied amplicon and metagenome sequencing to investigate the bacterial and fungal communities among four compartments (non-rhizosphere, rhizosphere, episphere, and endosphere) of mangrove roots. We found different distribution patterns for both bacterial and fungal communities in all four root compartments, which could be largely due to niche differentiation along the root compartments and exudation effects of mangrove roots. The functional pattern for bacterial and fungal communities was also divergent within the compartments. The endosphere harbored more genes involved in carbohydrate metabolism, lipid transport, and methane production, and fewer genes were found to be involved in sulfur reduction compared to other compartments. The dynamics of root-associated microbial communities revealed that 56–74% of endosphere bacterial taxa were derived from non-rhizosphere, whereas no fungal OTUs of non-rhizosphere were detected in the endosphere. This indicates that roots may play a more strictly selective role in the assembly of the fungal community compared to the endosphere bacterial community, which is consistent with the projections established in an amplification-selection model. This study reveals the divergence in the diversity and function of root-associated microbial communities along a continuous fine-scale niche, thereby highlighting a strictly selective role of soil-root interfaces in shaping the fungal community structure in the mangrove root systems.

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INTRODUCTION

Mangroves account for 60–70% of tropical and sub-tropical coastlines worldwide and have tremendous ecological importance as they participate in elemental cycling, mediate global climate change, protect coastlines, and facilitate phytoremediation. Similar to typical terrestrial plants, mangroves depend upon mutually beneficial interactions with microbial communities. In particular, microbes residing in developed roots could help mangroves transform nutrients into usable forms prior to plant assimilation. These microbes also provide mangrove phyto-hormones for suppressing phytopathogens or helping mangroves withstand heat and salinity. In turn, root-associated microbes receive carbon metabolites from the plant via root exudates, thus close associations between the plant and microbes are established for their mutual benefits.

Highly diverse microbial communities (mainly bacteria and fungi) have been found to inhabit and function in mangrove roots. For example, diazotrophic bacteria in the vicinity of mangrove roots could perform biological nitrogen fixation, which provides 40–60% of the total nitrogen required by mangroves. The soil attached to mangrove roots lacks oxygen but is rich in organic matter, providing an optimal microenvironment for sulfate-reducing bacteria (SRB) and methanogens. Ligninolytic, cellulolytic, and amylolytic fungi are prevalent in the mangrove root environment. Rhizosphere fungi could help mangroves survive in waterlogged and nutrient-restricted environments. These studies have provided increasing evidences to support the importance of root-associated bacteria and fungi for mangrove growth and health. However, systematic field studies on the overall taxonomic and functional diversity of mangrove root-associated microbial communities are still limited beyond examples of specific types of functional microbial members.

Recent studies have investigated the detailed structure of root-associated microbial communities at a continuous fine-scale in other plants, where a microhabitat was divided into four root compartments: endosphere, episphere, rhizosphere, and non-rhizosphere. Moreover, the microbial communities in each compartment have been reported to have unique characteristics. The rhizosphere could emit root exudates that selectively enriched specific microbial populations; however, these exudates were found to exert only marginal impacts on microbes in the non-rhizosphere soil. Furthermore, it was noted that the root episphere, rather than the rhizosphere, was primarily responsible for controlling the entry of specific microbial populations into the root, resulting in the selective enrichment of Proteobacteria in the endosphere. These findings provide new insights into the niche differentiation of root-associated microbial communities. Nevertheless, amplicon-based community profiling may not provide the functional characteristics of root-associated microbial communities in plant growth and biogeochemical cycling. Unraveling functional patterns across the four root compartments holds a great potential for understanding functional mechanisms responsible for mediating...
root–microbe interactions in support of enhancing mangrove ecosystem functioning.

Recently, root exudates were reported to be well-known determinants of root-associated microbial assemblages. Root exudation is a spatially defined process that contributes to distinct microbial communities that have been linked to specific root compartments. This is partly because root exudates could act as a carbon source and alter the rhizosphere pH (24, 15). However, the effect of root exudates on rhizobiome assembly is complicated, and the strategy of root-associated microbial community assembly at the soil-root interface remains controversial. The findings of some studies on root-associated microbiota in rice corroborate a two-step or multiple-step model in the root microbiota assembly, where specific microbial taxa in soil gradually become depleted or enriched during root colonization (22, 25). Other studies on rice and Medicago root-associated microbiota pointed to the applicability of the amplification-selection process, where dominant phyla would undergo substantial enrichment in the rhizosphere followed by the specific recruitment of certain phyla into the roots (23).

We aimed to determine the assembly processes that shape mangrove root-associated microbiota and examine the extent of differentiation and enrichment of mangrove root-specific microbial taxa across four root compartments. Furthermore, a community assembly framework developed by Vellend (24) and modified by Stegen et al. (25) allowed us to disentangle the ecological processes (heterogeneous selection, homogeneous selection, homogeneous dispersal, dispersal limitation, and undominated processes) that drive the mangrove root-associated microbial community composition at the spatial scale.

In this study, we used 16S rRNA and internal transcribed spacer (ITS) gene amplicon and metagenomic sequencing to systematically explore the diversity, function, and assembly of root-associated microbial communities in four continuous fine-scale root compartments (nonrhizosphere, rhizosphere, episphere, and endosphere) of Kandelia obovata (KO), a native mangrove plant of Southern China (24). We hypothesized that the diversity and function of mangrove root-associated microbial communities at the finescale could be largely affected by niche differentiation along the root compartments, and that the secretion of root exudates would affect root microbiomes in microbe-soil-plant systems (15). This study provides new insights into the understanding of root-associated microbial communities and their assembly mechanisms in mangrove ecosystems.

RESULTS

Diversity and composition of microbial communities among four mangrove root compartments

To determine whether microbial diversity varied across four continuous fine-scale compartments, we analyzed mangrove root-associated bacterial and fungal communities by sequencing 16S rRNA and ITS gene amplicons. Our amplicon sequencing analysis revealed a substantial difference in the diversity of bacterial and fungal communities among the compartments of nonrhizosphere (N), rhizosphere (R), episphere (P), and endosphere (D) (Fig. 1a). We observed the lowest Shannon-diversity and operational taxonomic unit (OTU) richness in the endosphere for both bacterial and fungal communities. The non-rhizosphere exhibited a higher bacterial diversity than the episphere, although fungal diversity indices were similar among the three exterior compartments (N, R, and P). Principal coordinate analysis (PCoA) based on Bray–Curtis distances showed that bacterial and fungal communities in the four root compartments were well-separated (P < 0.05, Adonis test), with the endosphere samples distinctly separated from the samples in other compartments (Fig. 1b).

The detected OTUs were distributed across seven dominant phyla. The abundance of Proteobacteria increased gradually from the non-rhizosphere to the endosphere, whereas the abundance of Chloroflexi deceased (Supplementary Fig. 1). As the most distinctive compartment, the endosphere was dominated by Proteobacteria (69.83%), and contained a low abundance of Chloroflexi (7.93%) and Actinobacteria (4.24%) (Supplementary Fig. 1a). The enrichment of a highly diverse Proteobacteria community in the endosphere was accompanied by the dominance of the following bacterial families (Supplementary Fig. 1b): two Alpha-proteobacterial families (Hyphomicrobiaceae and Rhodobacteraceae, 11.25% and 5.24%, respectively), two Gamma-proteobacterial families (Vibrionaceae and Saccharospirillaceae, 11.48% and 2.94%, respectively), and one Delta-proteobacterial family (Desulfovibrillaceae, 10.72%). At the genus level, we also observed notable differences among the four root compartments. The endosphere had a significantly greater proportion of Vibrio and Saccharospirillum than other compartments (P < 0.05, Student’s t-test), whereas Desulfooccus, Desulfococcaceae, and Delfuivitaleae were mostly depleted in the endosphere compared to the other three compartments (Supplementary Fig. 1c). In line with bacterial communities, fungal communities also showed significant variations across the four root compartments (P < 0.05, Student’s t-test). As the two dominant known fungal phyla in the four root compartments, Ascomycota (11%) and Basidiomycota (17%) in the endosphere had a lower abundance than those in other compartments (Ascomycota: 26%; Basidiomycota: 22%) (Supplementary Fig. 2). This variation trend across the four root compartments was also apparent in the abundant OTUs belonging to the unclassified fungi (Supplementary Fig. 2).

Enriched or depleted microbial OTUs among four mangrove root compartments

To identify the OTUs that contributed to the divergence in microbial community composition in the four root compartments, we conducted differential abundance analyses using a negative binomial distribution with OTU counts. Non-rhizosphere soils were set as the control and 0.01 as the adjusted P value cutoff. From the exterior (rhizosphere) to interior (endosphere) root compartments, the number of detected OTUs in bacterial communities was similar (R-P-D: 1133-1258-1239), but that in fungal communities showed a decreasing trend (R-P-D: 1163-934-732). The numbers of the enriched bacterial OTUs (R-P-D: 466-592-975) and the depleted fungal OTUs (R-P-D: 514-472-246) exhibited a completely different trend. Such differences were notable in the endosphere, which harbored more depleted bacterial OTUs (975) but less enriched bacterial OTUs (264) and fungal OTUs (246) compared with those in the episphere and rhizosphere (Fig. 1c).

We also observed noteworthy overlaps and distinctions in the abundant OTUs in the four root compartments. We found that 73.99% and 26.1% of rhizosphere-enriched bacterial OTUs were enriched in the episphere and endosphere, respectively. Likewise, 53.7% and 27.0% of rhizosphere-enriched fungal OTUs were also enriched in the episphere and endosphere, respectively. These data suggest that many microbes in the rhizosphere were able to colonize the root. Apart from these overlaps, considerable distinct OTUs were observed in each root compartment, as almost all of the depleted OTUs in the rhizosphere were depleted in the episphere, endosphere or both. For example, 55.2% of bacterial OTUs and 77.8% of fungal OTUs depleted in the episphere were also significantly depleted in the endosphere (P < 0.05) (Fig. 1d). Collectively, root-associated bacterial and fungal communities formed four spatially separable root compartments with distinct and overlapping microbial taxa.

Functions of root-associated microbial communities in the four mangrove root compartments

To explore microbial functions in the four root compartments, we analyzed mangrove root-associated microbial communities using shotgun metagenome sequencing with a focus on the relative abundance of key functional genes and pathways involved in...
carbon, nitrogen, sulfur and methane cycling (Fig. 2). The metagenomic contigs were annotated using evolutionary genealogy of genes: Non-supervised Orthologous Groups (eggNOG), Carbohydrate-Active enZymes (CAZy), and Kyoto Encyclopedia of Genes and Genomes (KEGG) databases. We found that the niche differentiation of root-associated microbial communities was accompanied by a notable divergence of microbial functions among the four root compartments (Fig. 2).

First, the functional annotations by eggNOG process categories indicated that the genes involved in carbohydrate transport and metabolism (G) were more abundant in the endosphere than in the rhizosphere and episphere (Fig. 2a). Also, the genes involved in secondary metabolite biosynthesis, transport, and catabolism (Q) were enriched in the endosphere and episphere, while they were depleted in the rhizosphere and non-rhizosphere. The endosphere, episphere, and rhizosphere consisted of many genes for lipid transport and metabolism (I) and signal transduction mechanisms (T). Through the analysis of CAZy reference sequences that are closely related to carbon cycling (Fig. 2b), we observed that the endosphere had the highest abundance of six CAZy families, with the enrichment of carbohydrate esterase (CE), glucosyltransferase (GT), and glycoside hydrolase (GH) genes (Fig. 2b).

Second, as nitrogen limitation of mangrove sediments can restrict plant growth and microbial activity, we estimated the abundance of a variety of functional genes involved in nitrogen cycling in the mangrove root environment (Fig. 2c). We found an increase in the relative abundance of genes involved in the conversion of extracellular polymers into NO$_3^-$ ($nrtA/B/C$) from the outside to the inside of root [gene transcripts per million (TPM) values in N and D were 1572 and 2986, respectively]. Although genes that allow the conversion of NO$_2^-$, NH$_3$OH, and N$_2$ ($nirB$, $hcp$, and $nifK$, respectively) were enriched in the four root compartments (Fig. 2c), the endosphere had the highest abundance of $nifK$ (934) and $nirB$ (2119) and a lower abundance of $nirS$ compared to other compartments (Fig. 2c).

Also, for sulfur cycling genes, the average abundance of $soxB/A/C/Z/Y/X/D$ in soil (7202) was higher than that in the rhizosphere, episphere, and endosphere (R-P-D: 6702-6623-6651), indicating that the potential of S$_2$O$_3^{2-}$ conversion to SO$_4^{2-}$ decreased around the root. Functional genes, including $sat$, $cysD$, $phsA$, and $dssB$, showed the lowest abundance in the endosphere, whereas $dssA$ was of minimum abundance in the episphere (Fig. 2d). These data indicate that sulfur reduction potential may be greater in the non-rhizosphere and rhizosphere soils.

In addition, mangrove ecosystems have been proposed as important methane sinks, thus we estimated the abundance of functional genes related to methanogenesis and methanotrophy (Fig. 2e). Our data indicated that the abundance of methyl coenzyme M reductase gene ($mcrA$) in the non-rhizosphere (17) was lower than that in the endosphere (69), where the particulate methane monoxygenase ($pmoA$) was of the lowest abundance.
The highest ratio of \( \text{mcrA} \) to \( \text{pmoA} \) (27) was observed in the endosphere, suggesting that mangrove roots have a high potential for CH\(_4\) production. Altogether, the above results revealed that root-associated microbial communities could play an important role in carbon, nitrogen, and sulfur cycling. More importantly, metabolic functional potentials differed substantially among the four mangrove root compartments.

Root exudates could shape mangrove root-associated microbial communities

To test whether root exudates regulate the diversity and composition of mangrove root-associated microbial communities, we measured the components of \( K. \) obovata root exudates using untargeted metabolomic analysis with negative mode acquisition (NEG) and positive mode acquisition (POS) modules. We identified 216 metabolites in mangrove root exudates, including amino acids, organic acids, polyhydroxy acids, sugars, phosphates, polyols, and N-compounds (Fig. 3a). The most abundant metabolites identified in the NEG module included palmitic acid (34.7%), stearic acid (16.4%), dehydroabietic acid (7.1%), oleic acid (6.7%), and myristic acid (6.1%), and those identified in POS module were dioctyl phthalate (43.7%), betaine (16.9%), phthalic acid mono-2-ethylhexyl ester (11.4%), and ethyl 3-hydroxybutyrate (3.5%).

Actinobacteria, Nitrospirae (diazotrophs), and mycorrhizal fungi, as members of root-associated microbial communities, have been reportedly to be correlated with some fatty acids (palmitic, linoleic, oleic, and stearic)\(^8\); we, therefore, compared the relative abundance of these populations among the four root compartments. Our data showed that the relative abundance of Actinobacteria (7.4%) was prominently higher in the episphere than in other compartments (Fig. 3c), which corroborates the finding that Actinobacterial members were the major groups stimulated by plant root exudates\(^27\). Likewise, diazotrophs were also specifically enriched in the rhizosphere and episphere (Fig. 3d, e). Other microbial taxa associated with root exudates\(^8,9\) such as Firmicutes, Chloroflex, and Ascomycota, were also significantly enriched (\( P < 0.05 \)) in the episphere (Fig. 1e). Considering non-rhizosphere as the control, we found that \( H. \) sp. in the episphere and rhizosphere was positively correlated with a variety of abundant constituents of root exudates, including betaine, salicylic acid, myristic acid, oleic acid, stearic acid, and palmitic acid (Supplementary Fig. 3). Therefore, we proposed that root exudates in mangroves could profoundly influence many root-associated microbial populations, especially those inhabiting the episphere.

Assembly mechanisms for microbial communities in mangrove root compartments

To understand the dynamics of mangrove root-associated microbial communities, we combined the use of source-tracker and ecological process analyses to elucidate microbial acquisition along the soil-root continuum. The results revealed that most bacterial OTUs in the endosphere were derived from the non-rhizosphere (74%), and only a small part of the episphere bacterial communities (4%) were detected in the endosphere compartment (Fig. 4a). Moreover, no fungal OTUs that were previously detected in the non-rhizosphere or episphere were detected in the endosphere (Fig. 4b). To verify the dissimilar manner of acquisition of bacteria and fungi by the root, we used real-time quantitative PCR (qPCR) to determine the ratio of bacteria to fungi among the four root compartments.
Supplementary Fig. 4). The results showed that the ratio of bacteria to fungi decreased from 128.0% in the non-rhizosphere to 13.8% in the endosphere. Furthermore, ecological process analysis revealed that the deterministic processes of heterogeneous selection contributed >40% to the community assembly for both bacteria and fungi (Fig. 4c). Altogether, the results suggested that the episphere could effectively act as a gate for controlling the entry of microbes into the root endosphere, revealing a dominance of heterogeneous selection in the assembly of the mangrove root-associated microbial communities. This is most probably linked to the low alpha diversity detected in the endosphere communities (Fig. 1a), which were selectively recruited from the exterior compartment communities.

**DISCUSSION**

Understanding the diversity and function of microbial communities along a root-associated fine-scale is crucial in elucidating microbial assembly mechanisms and their ecological importance in mangrove ecosystems. In this study, we analyzed the microbial diversity and their functions across the four root compartments, and their relationships with root exudates. Our results generally support the core hypothesis that the diversity and function of mangrove root-associated microbial communities would diverge along such continuous fine-scale niches. The lower diversity of both bacterial and fungal communities in the endosphere of mangrove root than the diversity of communities in other compartments, was similar to that in the root microbiome of...
Arabidopsis\textsuperscript{28}. This points to the selective role of the rhizoplane in controlling microbial entry into the root and reducing microbial diversity\textsuperscript{28-30}. Notably, the microbial function of root-associated microbial communities showed certain patterns in different microenvironments or compartments\textsuperscript{2,6,15,31}. In previous studies, plant carbon sources and root exudates were reported to attract microbial populations specifically involved in carbon cycling to the rhizosphere or episphere\textsuperscript{8,32}. Carbon cycling requires a variety of carbohydrate-active enzymes\textsuperscript{33}, including GTs\textsuperscript{34}. Some GTs in wheat root environments could help resist pathogenic fungi\textsuperscript{35}, whereas other GTs had a self-detoxification mechanism\textsuperscript{36}. Also, some GTs could catalyze the activation of hormones in plants and improve the transport efficiency of sugars in roots\textsuperscript{34}. In line with these previous findings, our study showed that the abundance of GTs increased from non-rhizosphere, rhizosphere, and episphere to endosphere, suggesting that the internal root microenvironment may have high sugar conversion efficiency and self-detoxification potentials.

Both sediment and rhizosphere of mangrove ecosystems are rich in SRB\textsuperscript{10}. As vital decomposers of organic matter in anaerobic environments, SRB play a critical role in the mineralization of organic sulfur and production of available iron and phosphorus for other organisms in mangrove ecosystems\textsuperscript{12}. Similarly, we found that root-related compartments had abundant SRB (e.g., \textit{Desulfococcus} and \textit{Desulfosarcina}) and associated functional genes (e.g., \textit{phsA} and \textit{dsrA}). More strikingly, SRB and sulfur reduction-related genes were specifically enriched in the exterior compartments. This is probably due to the presence of oxygen and redox potential gradients in the rhizosphere\textsuperscript{37}, which may facilitate sulfate reduction in the non-rhizosphere and rhizosphere soils.

Mangrove ecosystems emit large amounts of methane\textsuperscript{3,38}. Previous studies have shown that CH\textsubscript{4} emissions were remarkably affected by the abundance of methanogens (identified by \textit{mcrA}) and methanotrophs (identified by \textit{pmoA})\textsuperscript{38,39}. In our study, the ratio of \textit{mcrA} to \textit{pmoA} was much higher in the endosphere than in other exterior compartments, indicating that the endosphere had a higher potential of CH\textsubscript{4} emissions than other compartments. In despite of the previous finding that mangrove sediment has been considered as a key methane sink, this study suggests that CH\textsubscript{4} production via mangrove roots, and then transport into the above-ground plant may be an alternative but previously unrecognized pathway in mangrove ecosystems, and this phenomenon has been described in previous studies on coastal saltmarsh of \textit{Phragmites}\textsuperscript{40}.

Mangrove ecosystems are characterized by rich organic carbon and hypersalinity\textsuperscript{5,10}. Under such conditions, ammoniated bacteria have a higher affinity than denitrifying bacteria, enabling dissimilatory nitrate reduction to ammonium (DNRA) to be the main pathway for preserving nitrogen\textsuperscript{33,41}. Consistently, genes related to DNRA processes were enriched in root-related compartments in this study. Loaded with the abundant diazotrophic microbes, the three inner compartments of mangrove roots were likely to be rich in ammonia nitrogen. The results indicate the divergence of microbial diversity, abundance, and functions in the four root compartments. Therefore, the endosphere had high sugar conversion efficiency and high potential for self-detoxification, CH\textsubscript{4} emission, and ammonia nitrogen reserve,
while sulfate reduction was stronger in the non-rhizosphere and rhizosphere than in the episphere or endosphere.

Root exudates can affect the microbial community composition in the root environment, especially in the episphere and rhizosphere microenvironments. In previous studies, the role of root exudates was reported to be the recruitment of various functional microbes to protect the host plant. The high abundance of phenolic acids in root exudates likely results in the expression of key genes involved in the production of antifungal substances. Also, root exudates can increase microbial biomass and diversity by providing a variety of organic matter, including fatty acids and sugars. In this study, we found that root exudates in mangroves include amino acids, organic acids, polyhydroxy acids, sugars, phosphates, polyols, and N-compounds. Among these, phenolic acids (e.g., phthalic acid, salicylic acid) and fatty acids (e.g., palmitic acid, myristic acid, and oleic acid) in high abundances could attract more probiotics that protect host plants. For example, we found that Hypromicrobium (a typical denitrifying bacterium), Nitrosopirene, and diazotrophs were positively correlated with the abundant root exudates in the episphere and rhizosphere of mangroves; the presence of these organisms may lead to fast nitrogen cycling in mangrove ecosystems as previously observed.

Deterministic and stochastic assembly occur simultaneously along successional chronosequences and drive the spatial distribution of microbial communities in many ecosystems. However, the assembly and selection preferences of root-associated microbial communities remain controversial. First, the relative abundance patterns of microbial communities in the root compartments follow certain rules of spatial variation. Similar patterns in rice and Arabidopsis microbiomes indicated that the endosphere had a higher proportion of Proteobacteria and Spirochetes than the rhizosphere or non-rhizosphere, whereas Acidobacteria and Chloroflexi were mostly depleted in the endosphere. In this study, our results also showed that the abundance of Proteobacteria gradually increased, whereas the abundance of Chloroflexi gradually decreased from the non-rhizosphere to the endosphere. Specifically, we found that SRB such as Desulfococcus, Desulfoarcina, and Defluvialatae, were mostly depleted in the endosphere than in the other three compartments, indicating that the relative abundance of microbial communities among the four root compartments of mangroves followed certain rules from non-rhizosphere to endosphere.

Second, the specific habitat characteristics of mangrove root microenvironments may have important impacts on their microbial community assembly. In previous studies, hydrologic connectivity was reported to be a major factor in structuring microbial communities in most ecosystems and stochasticty was dominant in the assembly of aquatic environments, including ground-water, bioreactor, and flooded rice paddies. On the contrary, some host-associated bacterial communities, including animal guts, reportedly had a higher proportion of deterministic processes than stochastic processes. In this study, we found that deterministic processes were dominant in the four mangrove root compartments, suggesting that the microbial colonization in mangrove roots was not a passive process and that mangrove plants had a strong selectivity for their associated communities. In line with this notion, Fan et al. observed a decreasing importance of deterministic processes in determining the diazotrophic communities in relation to distance from wheat roots.

Third, many studies indicate that root microbiome acquisition is a continuous process of gradual filtration (two-step or multiple-step selection process), whereas the amplification of microbes in the rhizosphere/episphere prior to the selection of microbes in the rhizosphere/episphere (amplification-selection process) was preferred in other studies. In the present study, many dominant microbial phyla (e.g., Proteobacteria, Spirochetes, Firmicutes, Acidobacteria, and Actinobacteria) were enriched in the episphere to certain degrees; specific phyla (e.g., Proteobacteria, Spirochetes, and Acidobacteria) were selectively enriched in the endosphere. These results revealed that the assembly pattern of mangrove root microbial communities could depend on amplification-selection. To the best of our knowledge, there are no studies wherein assembly processes combined with chronosequence with spatial patterns were conducted in mangrove ecosystems. It is crucial to further explore the factors that influence such assembly processes and underlying mechanisms.

In this study, we showed the spatial and exudation effects of mangrove roots on the diversity, function, and assembly of root-associated microbial communities. Root-associated microbiomes could form four spatially separable compartments and exhibit divergent diversity and function patterns in the mangrove root environment. Also, we found that root exudates played an important role in the development of root microbiome in the episphere and rhizosphere compartments. In addition, each mangrove root compartment had unique ecological niches for their bacterial and fungal communities. The assembly mechanisms appeared to be represented by the amplification-selection model. This study provides new insights into the understanding of microbial diversity, function, and their assembly mechanisms in the mangrove root environment. Future studies are needed to clarify the mechanisms by which root exudation affects the root microbiome and to explore the microbe-soil-plant interactions in mangrove ecosystems.

METHODS

Sampling sites, root collection, and environmental properties

The sampling site was located at Shuidong Bay of Maoming City (21°30′38.82″N, 111°0′37.27″E) (Supplementary Fig. 5), Guangdong, China, where natural mangrove communities are dominated by K. obtusata. Six individual KO saplings were extracted in April 2019, and their root-associated samples were fractionated into four compartments (Supplementary Fig. 6): non-rhizosphere soil (N), rhizosphere soil (R), episphere (P), and endosphere (E). The samples for these compartments were processed as described by Duran et al. and Edwards et al. Briefly, roots were collected from mangrove plants, and non-rhizosphere soil was separated from the root by shaking. The rhizosphere soil (1 mm thickness around the root) that could not be removed by shaking was collected by washing with sterile water. The clean roots were then washed three times to remove the remaining soil and placed into 1× TE buffer supplemented with 0.1% Triton X-100 in a 50 mL Falcon tube. Next, the episphere samples were collected via shaking and extensive shaking in 1× TE buffer supplemented with 0.1% Triton X-100. The episphere microbial biomass was collected via filtering the resulting suspension through 0.22μm pore size membranes (Nuclepore, Whatman, Meterstone, UK). To collect the endosphere microbial biomass, the roots were surface-sterilized for 1 min in 80% ethanol and subsequently sterilized again for 1 min in 0.25% NaClO. All four root compartment samples were stored at −80°C until DNA extraction. For non-rhizosphere soils, the moisture content, pH, salinity, oxidation reduction potential, ammonium-N, nitrate-N, nitrite-N, total carbon, and total nitrogen were measured as previously described.

DNA extraction, PCR amplification, and sequencing

Approximately 0.5 g non-rhizosphere and rhizosphere soil with six replicates was used for DNA extraction using a Power Soil DNA Isolation Kit (MoBio, Carlsbad, CA, USA) according to the manufacturer’s instructions with the modified sodium dodecyl sulfate extraction method. The episphere compartment DNA was extracted using a Power Water DNA Isolation Kit (MoBio, Carlsbad, CA, USA) according to the manufacturer’s instructions. DNA from endosphere samples was extracted using a Power Plant DNA Isolation Kit (Mo Bio Laboratories, Inc., Carlsbad, CA, USA) after thorough rinsing under liquid nitrogen. The DNA quality based on 260/280 and 260/230 nm ratios was assessed using Nano Drop ND-2000 Spectrophotometer (Thermo Fisher Scientific, MA, USA). DNA samples with good quality were diluted to 2 ng/μL for subsequent PCR amplification.
Sequence analysis of 16S rRNA and ITS1 gene amplicons

Raw sequences were first processed using Trimmomatic59 and FLASH60. During filtering, the sequences were trimmed with a moving window of 50-bp and a quality threshold score of 30. The dataset was then simplified by eliminating singletons. Paired 16S rRNA amplicon sequences were then clustered into OTUs by UPARSE61 based on a 97% sequence identity using Quantitative Insights into Microbial Ecology62 open reference OTU picking strategy with the Greengenes 16S rRNA database (v.13.5) as a reference63. The sequences matching “Chloroplast” and “Mitochondria” were excluded from the datasets. ITS sequences were processed using ITS464 and clustered at 97% sequence identity using UPARSE61. Fungal OTUs were checked for chimeric sequences using the Uchime reference against a dedicated chimera detection database65. We obtained a total of 63,590 and 63,712 high-quality 16S rRNA and ITS gene amplicon sequences per sample, respectively (Supplementary Table 1 and Supplementary Table 2). With >97% sequence identity and removal of low-abundance OTUs (<0.01% of total abundance), the 16S rRNA sequences were clustered into 12,664 OTUs, and ITS sequences into 15,241 OTUs for all samples (Supplementary Table 3).

Shotgun metagenome sequencing data analysis

One microgram of DNA was used for metagenome sequencing library preparations combined with NEBNext UltraTM DNA Library Prep Kit for Illumina (NEB, USA) according to the manufacturer’s recommendations. Index codes were added to attribute sequences to each sample; these preparations combined with NEBNext UltraTM DNA Library Prep Kit for Illumina (NEB, USA) according to the manufacturer’s recommendations. Index codes were added to attribute sequences to each sample; these preparations were purifed (AMPure XP system), and the libraries were checked (Bioanalyzer). The resulting solution was then centrifuged at 22,542 × g for 15 min at 4 °C. The 400 μL supernatant was transferred to a sterile tube and dried in a vacuum concentrator at 37 °C. Thereafter, the dried sample was reconstituted in 200 μL 50% acetonitrile via sonication on ice for 10 min. The resulting solution was then centrifuged at 22,542 × g for 15 min at 4 °C. The 75 μL supernatant was transferred to sterile glass vial for liquid chromatography–mass spectrometry (LC/MS) analysis. LC-MS/MS analysis was performed using 1290 Infinity series UHPLC System (Agilent Technologies) equipped with a UPLC BEH Amide column (2.1 × 100 mm, 1.7 μm, Waters). Ion spray voltage floating (ISVF) at 5000 V or −4000 V was applied in the positive or negative modes, respectively. MS raw data files were converted to the mzXML format using ProteoWizard, and processed using the R package XCMS (version 3.2). The data analysis includes peak deconvolution, alignment, and integration processes. The minfrac and cutoff were set as 0.5 and 0.3, respectively. The in-house MS2 database was applied for metabolite identification.

Statistical analysis

Statistical analyses were performed using the VEGAN package75 in R 3.6.0, including alpha-diversity indices (Shannon index and Chao index) and PCOA. The Student’s t-test was performed using SPS5. Applied to test the significant differences in microbial abundance among compartments. Differentially abundant OTUs were detected using Deseq2 generalized linear model approach76.

Source-Tracker analysis, a Bayesian approach, was used to identify microbial communities in an environmental sink to various potential sources77. SourceTracker analysis was conducted within an in-house pipeline (http://mem.reesc.ac.8080) which consisted of relevant bioinformatics tools. The percentage value was the statistical average of the Source-Tracker results. For the ecological process analysis, we calculated the βMNTD (β-mean-nearest taxon distance) using the R function “comdist” for the phylogenetic distance between each OTU in one community (k) and its closest relative in a second community (m). We used βNTI in combination with Bray–Curtis-based Raup–Crick (RCpy) to quantify the contribution of major ecological processes to the assembly of root-associated microbial communities67,80. The percentage value was derived from the statistical average of the ecological process results.

Reporting summary

Further information on research design is available in the Nature Research Reporting Summary linked to this article.
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