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Fungal community remediate quartz tailings soil under plant combined with urban sludge treatments

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Introduction: Tailings can cause extensive damage to soil structure and microbial community. Phytoremediation is an effective strategy for remedied tailings soil due to its environmentally friendly and low-cost advantage. Fungi play a crucial role in nutrient cycling, stress resistance, stabilizing soil structure, and promoting plant growth. However, the fungal community variation in phytoremediation remains largely unexplored.

Methods: We analyzed soil fungal community based on high-throughput sequencing during three plant species combined with urban sludge to remediate quartz tailings soil.

Results: The results indicated that the fungal diversity was significantly increased with plant diversity, and the highest fungal diversity was in the three plant species combination treatments. Moreover, the fungal diversity was significantly decreased with the addition of urban sludge compared with plant treatments, while the abundance of potential beneficial fungi such as *Cutaneotrichosporon*, *Apiotrichum*, and *Alternaria* were increased. Notably, the fungal community composition in different plant species combination treatments were significant difference at the genus level. The addition of urban sludge increased pH, available phosphorus (AP), and available nitrogen (AN) content that were the main drivers for fungal community composition. Furthermore, the fungal networks of the plant treatments had more nodes and edges, higher connectedness, and lower modularity than plant combined with urban sludge treatments.

Conclusion: Our results showed that three plant species combined with urban sludge treatments improved fungal community and soil properties. Our results provide insights for quartz tailings soil remediation using plant-fungi-urban sludge.

KEYWORDS

fungal diversity, fungal community structure, phytoremediation, quartz tailings, phosphorus and nitrogen, urban sludge

1. Introduction

The exploitation of mineral resources that play an essential role in the development of humans was shown exponential growth as the economy development (Carvalho, 2017). However, the vast amount of waste liquids and rocks from the mining process are transported into soil and then destroy the ecological environment (Kossoff et al., 2014; Adiansyah et al., 2015). It is difficult for plants to grow on tailings soil due to their physicochemical characteristics, such as lower or higher pH, higher heavy metal pollution, lower water retention capacity, deficiencies in soil organic matter, and lack of plant growth nutrients (Sheoran et al., 2008; Wang et al., 2017; da Silva et al., 2022). Notably, the changes in soil physicochemical properties can affect the soil microbial community structure (Jia et al., 2019). The common restoration methods for tailings soil are biological, physical, and chemical remediation (Festin et al., 2018). Phytoremediation including plants and related microorganisms that have the advantages of low-cost, sustainable, and environmentally friendly technology, are ideally used to remediate tailings soil compared with physical and chemical remediation (Dickinson, 2017; Sharma, 2020).

More than 400 kinds of plant species in the world can metabolize pollutants including accumulating organic and inorganic contaminants by improving microbial degradation of contaminants in the root zone (Arthur et al., 2005; Faucon et al., 2007). Furthermore, plants can transport, accumulate or degrade soil heavy metal pollutions such as Zn, Pb, Cd, Mn, Cu, Cr, Fe, As, and Ni, and then reduce environmental impact through litter decomposition, root exudates and soil properties (Wang et al., 2015; Sanchez et al., 2018; Ali et al., 2020; Su et al., 2022). For example, some species of Legumes are often used to improve the nitrogen content of tailings soils (Young et al., 2015). Moreover, some plant species can tolerate drought of semiarid and mining soil with low water holding capacity (Santibañez et al., 2012). Therefore, suitable plant species are very important to repair tailings soil.

Fungal community are critical factors in maintaining plant biodiversity, soil health and productivity, and soil biogeochemical processes such as nitrogen and phosphorus cycling (Mangan et al., 2010; Urbanová et al., 2015). They contribute to plant disease control and growth promotion, and plant stress resistance (Ozimek and Hanaka, 2020). Some fungal groups, such as *Penicillium*, can assist plant in phosphorus and nitrogen uptake in heavy metal-contaminated soil (Ikram et al., 2018; Elfiati et al., 2021). Moreover, some species of Basidiomycetes can improve plant salt tolerance by increasing the concentration of osmotic fluid in plant cells, and enhancing plant minerals uptake and potassium ions for metal-detoxifying (Bukhori et al., 2020). Notably, arbuscular mycorrhizal fungi (AMF) can assist plant to facilitate mineral and water uptake (Debeljak et al., 2018; Begum et al., 2019). *Archaeorhizomyces* enhance stress resistance and inhibit disease by increasing the bioactive components of the plant (Zhang et al., 2020). Furthermore, the plant growth promoting fungi (PGPR) have been widely used to prevent and control heavy metal pollution due to its ability to promote plant growth and induce host resistance (Geetha et al., 2022). Moreover, fungi also play a crucial role in maintaining soil plant productivity (Frac et al., 2018). These findings suggest that fungal community play an essential role in

plant defense against stressful environment and facilitate nutrient absorption. In turn, plant can significantly improve the diversity and abundance of soil microbial community by providing an ideal environment to inhabit (Zhao A. et al., 2019; Wu et al., 2022). Plant can recruit beneficial fungal community through root systems, stems, leaves, and even seeds (Santoyo, 2021). Higher microbial diversity led to higher microbial function level and then accelerate tailings soil remediation (Fuke et al., 2021). However, plant recruit which fungal community and play what roles in tailings soil remediation remain unclear.

Extreme environmental conditions strongly influence the efficiency of phytoremediation (Chandra and Kumar, 2018). In order to improve the recovery efficiency, some amendment combination with plants have been used in the rehabilitation of tailings soil (Asensio et al., 2013b; Alcantara et al., 2015). Notably, some studies have shown that urban sludge significantly neutralized soil pH in restoring Cu tailings (Asensio et al., 2013a). In addition, urban sludge can improve soil properties and influence soil bacterial community structure (Bai et al., 2019; Zuo et al., 2019). Significantly, urban sludge is often used as an amendment to phytoremediation due to its easy availability and rich in inorganic and organic nutrients. However, we still need to better understand how the plant species and urban sludge improve the properties and biological condition of tailings soil. In this study, we used three plant species (*Lolium perenne* L., *Vicia sepium* L., and *Medicago sativa* L.) combined with urban sludge to remediate quartz tailings soil. We analyzed the variation and co-occurrence of fungal community in 14 treatments. We hypothesized that (i) the treatments of plant species combination could significantly increase fungal diversity than single species. (ii) Plant could recruit beneficial fungi to resist stress in tailings soil. (iii) Plant combined with urban sludge could effectively remediate tailings soil due to the improvement of biological-chemical-physical properties. The results provide systematic information for the remediation of tailings soil and fully reveal the relationship among plant, urban sludge, and soil fungal community.

2. Materials and methods

2.1. Material preparation

Quartz tailings soil and urban sludge were collected separately from the Fengyang County mining area and Chuzhou Zhongye Huatian Water Co., Ltd. of Chuzhou City, Anhui Province, China. Impurities on the surface of quartz tailings soil were removed during sampling, and quartz tailings soil were randomly collected and brought back to the laboratory for remediation test. Three soil samples were spread in a ventilated place to dry, and then passed through 2 mm sieve to remove impurities. The sieved soil samples were used to measure soil physical and chemical properties. The urban sludge reached the discharge standard after being treated by the providing company, and the properties such as nutrients and harmful substances meet the Chinese national standards.

Vicia sepium L.(Y) and *M. sativa* L.(Z) belong to Legume that have the characteristic of drought and barren tolerance, salt and alkali resistance, and strong ability to enrich heavy metals (Templeton, 2018; Bao et al., 2022). *Lolium perenne* L.(H) is widely

used in tailings soil due to its strong resistance and growth ability (Irhema, 2019). We used the three plant species to remediate tailings soil. The seeds of the three plant species were purchased from Century Tianyuan (Luoyang) Ecological Technology Co., Ltd. (Henan Province, China). Seeds were soaked in sterile water for 12 h at room temperature, then loaded into sterilized Petri dishes, and covered with wet gauze until germination. Seedlings that were strong and uniform in size were selected for the subsequent treatment.

2.2. Experimental design

The experiment was conducted in a greenhouse using potted plants placed in polyethylene plastic pots (upper diameter: 16 cm, lower diameter: 12 cm, height: 17 cm, volume: 2 L). Approximately 1 kg of quartz tailing soil was placed in each pot. Quartz tailings soil was treated with plant or plant combined with urban sludge, respectively. The treatments of plant species were divided into one, two and three plant species combination. Therefore, seven treatments were set in the plant treatment group, including Y treatment (KY), H treatment (KH), Z treatment (KZ), Y & H treatment (KYH), Y & Z treatment (KYZ), H & Z treatment (KHZ), and Y & H & Z treatment (KYHZ), respectively, (Figure 1A). Seven plant species combined with urban sludge treatments were set in Figure 1B. Two hundred gram of urban sludge were drizzled in the quartz tailing soil in each pot. Seven treatments in plant combined with urban sludge group were Y treatment (WY), H treatment (WH), Z treatment (WZ), Y & H treatment (WYH), Y & Z treatment (WYZ), H & Z treatment (WHZ), and Y & H & Z treatment (WYHZ), respectively. Six uniform seedlings were planted per pot. Three or two seedlings of each plant species in two plants and three plants mixed treatments were planted per pot, respectively. Each treatment was replicated five times. Watering was in demand during 100 days of treatment.

2.3. Quartz tailings soil properties and plant biomass analysis

Soil of 14 treatments were collected and sieved through a 2 mm sieve after the experiment, and then divided into two parts for soil properties determination and DNA extraction. The content of total nitrogen (TN), available phosphorus (AP) and nitrogen (AN), iron trioxide (Fe_2O_3), pH, and silicon dioxide (SiO_2) were measured according to the method of previous studies (Sun et al., 2015; Qi et al., 2017; Yu et al., 2017; Zhao S. et al., 2019). Briefly, pH was determined by pH meter (S500-F, Mettler Toledo, Germany) at a ratio of 1:2.5 (weight/volume) for soil versus distilled water. The content of TN was determined by the Kjeldahl digestion method. AP was extracted using 0.5 M NaHCO_3 and determined by molybdenum blue method. Alkaline hydrolysis diffusion was used to determine the soil AN. The content of Fe_2O_3 in soil was determined by atomic absorption spectroscopy. The content of SiO_2 was determined by UV-visible spectrophotometry. Furthermore, plants of each pot were harvested, and measured the aboveground and belowground biomass of each plant.

2.4. DNA extraction and sequencing

DNA of soil samples were extracted using the E. Z.N.ATM Mag-Bind Soil DNA Kit (Omega, United States). DNA quality was checked using 1% agarose gel electrophoresis and stored at -80°C for PCR amplification. The fungal ITS1 region was amplified using the primer ITS1F (5'-CTGGTCATTAGAGGAAGTAA-3') and ITS2R (5'-GCTGCGTTCCTCATCGATGC-3') on ABI GeneAmp 9,700 PCR thermocycler (ABI, CA, United States). The PCR products were recovered by cutting the gel using the AxyPrep DNA Gel Recovery Kit (Axygen Biosciences, United States), and then detection and quantification were performed with the QuantiFluorTM-ST blue fluorescence quantitative system (Promega, United States). Paired-end sequencing were performed by Majorbio Bio-Pharm Technology Co. Ltd. (Shanghai, China) on the Illumina MiSeq PE300 platform (Illumina, San Diego, CA, United States). All raw reads were deposited into NCBI Sequence Read Archive (SRA) database (Accession number: PRJNA923135). The raw sequences that contained ambiguous nucleotides, short length, and low quality were discarded (Sun et al., 2021). The high-quality sequences were analyzed by the QIIME (Caporaso et al., 2010). Operational taxonomic units (OTUs) at 97% similarity were identified using the UPARSE (Edgar, 2013). All sample sequences were flattened according to the minimum sequence number. 21,160 OTUs were acquired, and flattened at the OTUs level. OTUs scale generated after flattened were used for subsequent analysis.

2.5. Bioinformatics and statistical analysis

Alpha diversity indicators including Chao, and Shannon index were calculated using the vegan package in R at the OTUs level. The difference among treatments were analyzed using Student's *t*-test significant difference (Kembel et al., 2010). Fungal community composition at the phylum and genus level were generated using the "ggplot2" package (Villanueva and Chen, 2019). The difference of fungal community composition were evaluated using principal coordinate analysis (PCoA) based on the Bray-Curtis distance (Edgar, 2013). The relationship between fungal community and soil properties were analyzed based on Redundancy analysis (RDA). The relationship between soil properties and fungal diversity were performed using Pearson's correlation analyses (the top 20 at the genus level) (Capblancq and Forester, 2021; Cheng et al., 2021). Significant difference of the relative abundance of fungal community among different treatments were analyzed using the linear discriminant analysis effect size (LEfSe) (Chang et al., 2022). The LDA threshold was chosen four and used All-against-all to evaluate the statistical differences. The variation of fungal community composition was analyzed by multiple group comparisons based on Welch's *t*-test. The difference between plant biomass and soil properties were analyzed using the least significant difference (LSD) (Urbanová et al., 2015).

2.6. Network construction

The fungal network of plant treatments (PT) and plant combined with urban sludge treatments (PUT) were constructed

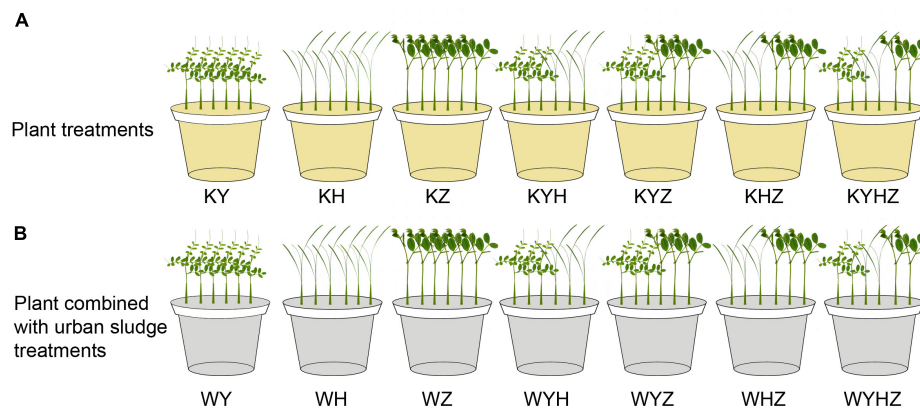


FIGURE 1

Experimental design diagram of quartz tailings soil remediation by plant and plant combined with urban sludge. Panel (A) represents plant treatments. Panel (B) represents plant combined with urban sludge treatments. Y, H, and Z represent *Vicia sepium* L., *Lolium perenne* L., and *Medicago sativa* L., respectively.

TABLE 1 Effect of plant and plant combined with urban sludge treatments on quartz tailings soil properties.

Treatment	Available nitrogen (AN, mg/kg)	Available phosphorus (AP, mg/kg)	Iron trioxide (Fe ₂ O ₃ , %)	Silicon dioxide (SiO ₂ , %)	Total carbon (TC, %)	pH
Untreated soil	14.00 ± 2.13 ^d	1.86 ± 0.15 ^{bc}	1.25 ± 0.13 ^d	52.83 ± 0.58 ^{abc}	0	5.35 ± 0.21 ^f
KY	10.62 ± 3.26 ^d	1.93 ± 0.09 ^c	1.82 ± 0.16 ^{bcd}	51.46 ± 0.42 ^{bc}	0	5.52 ± 0.06 ^{ef}
KH	11.18 ± 2.08 ^d	1.53 ± 0.05 ^c	2.40 ± 0.26 ^{ab}	52.79 ± 0.80 ^{abc}	0	7.14 ± 0.32 ^a
KZ	16.16 ± 1.79 ^d	1.49 ± 0.08 ^c	2.58 ± 0.06 ^a	53.28 ± 1.15 ^{ab}	0	6.58 ± 0.07 ^{bc}
WY	35.64 ± 5.15 ^{ab}	15.72 ± 1.29 ^b	1.91 ± 0.09 ^{bcd}	51.37 ± 1.20 ^{bc}	0	5.44 ± 0.07 ^f
WH	36.90 ± 4.12 ^a	39.42 ± 3.53 ^a	2.04 ± 0.28 ^{abc}	50.90 ± 0.72 ^c	0	6.42 ± 0.39 ^{bcd}
WZ	43.53 ± 7.35 ^a	40.17 ± 11.16 ^a	2.00 ± 0.24 ^{abc}	52.49 ± 0.90 ^{abc}	0	6.00 ± 0.03 ^{de}
KYH	12.59 ± 1.66 ^d	1.41 ± 0.22 ^c	2.57 ± 0.09 ^a	53.08 ± 0.87 ^{abc}	0	6.47 ± 0.13 ^{bcd}
KYZ	21.53 ± 5.07 ^{bcd}	0.79 ± 0.33 ^c	2.53 ± 0.09 ^a	52.47 ± 0.96 ^{abc}	0	6.50 ± 0.10 ^{bcd}
KHZ	19.43 ± 4.20 ^{cd}	0.58 ± 0.28 ^c	2.35 ± 0.15 ^{ab}	53.30 ± 1.14 ^{ab}	0	6.85 ± 0.22 ^{ab}
WYH	15.32 ± 3.12 ^d	8.78 ± 1.26 ^{bc}	1.83 ± 0.23 ^{bcd}	50.85 ± 0.60 ^c	0	6.19 ± 0.11 ^{cd}
WYZ	21.23 ± 0.78 ^{bcd}	13.79 ± 0.60 ^{bc}	1.74 ± 0.13 ^{cd}	53.96 ± 0.86 ^a	0	6.35 ± 0.09 ^{bcd}
WHZ	31.85 ± 4.69 ^{abc}	43.49 ± 10.13 ^a	1.88 ± 0.39 ^{bcd}	52.69 ± 0.71 ^{abc}	0	7.15 ± 0.30 ^a
KYHZ	13.56 ± 2.33 ^d	1.55 ± 0.04 ^c	2.30 ± 0.16 ^{abc}	52.16 ± 0.43 ^{abc}	0	6.43 ± 0.05 ^{bcd}
WYHZ	42.55 ± 13.47 ^a	38.96 ± 8.53 ^a	2.37 ± 0.34 ^{ab}	52.42 ± 0.39 ^{abc}	0	6.49 ± 0.22 ^{bcd}
Urban sludge	4111.33 ± 109.11	14.61 ± 0.38	3.25 ± 0.04	52.27 ± 0.41	0	6.03 ± 0.01

a, b, c, d, e, and f indicated significant differences ($P < 0.05$).

on integrated network analysis based on random matrix theory (RMT) on iNAP platform (Feng K. et al., 2022). The OTUs level with abundance of >0.01 were filled based on Spearman's rank correlation. Before the analysis, only OTUs detected in $>20\%$ of all samples were used for network construction. The largest and smallest cutoff value for RMT scanning was 1.0 and 0.01 separately. The module orders for each species were obtained by the fast-greedy method based on Spearman's correlation. The nodes of fungi were assigned to the peripheral, connector, module hub, or network hub, according to their patterns of within- and among-module connections (Z_i and P_i). The calculation was permuted by 100 times at each step of node removal at the proportion. Network properties, including node and edge number, connectedness, modularity, average path distance, and proportions of positive and negative

edges were calculated and selected for comparison of networks. Module-Eigen Gene analysis was used to analyze the relationship between modules and environmental factors. The globe network of PT and PUT were visualized by Gephi 0.9.2 with the "Fruchterman Reingold" layout algorithm (Bastian et al., 2019).

3. Results

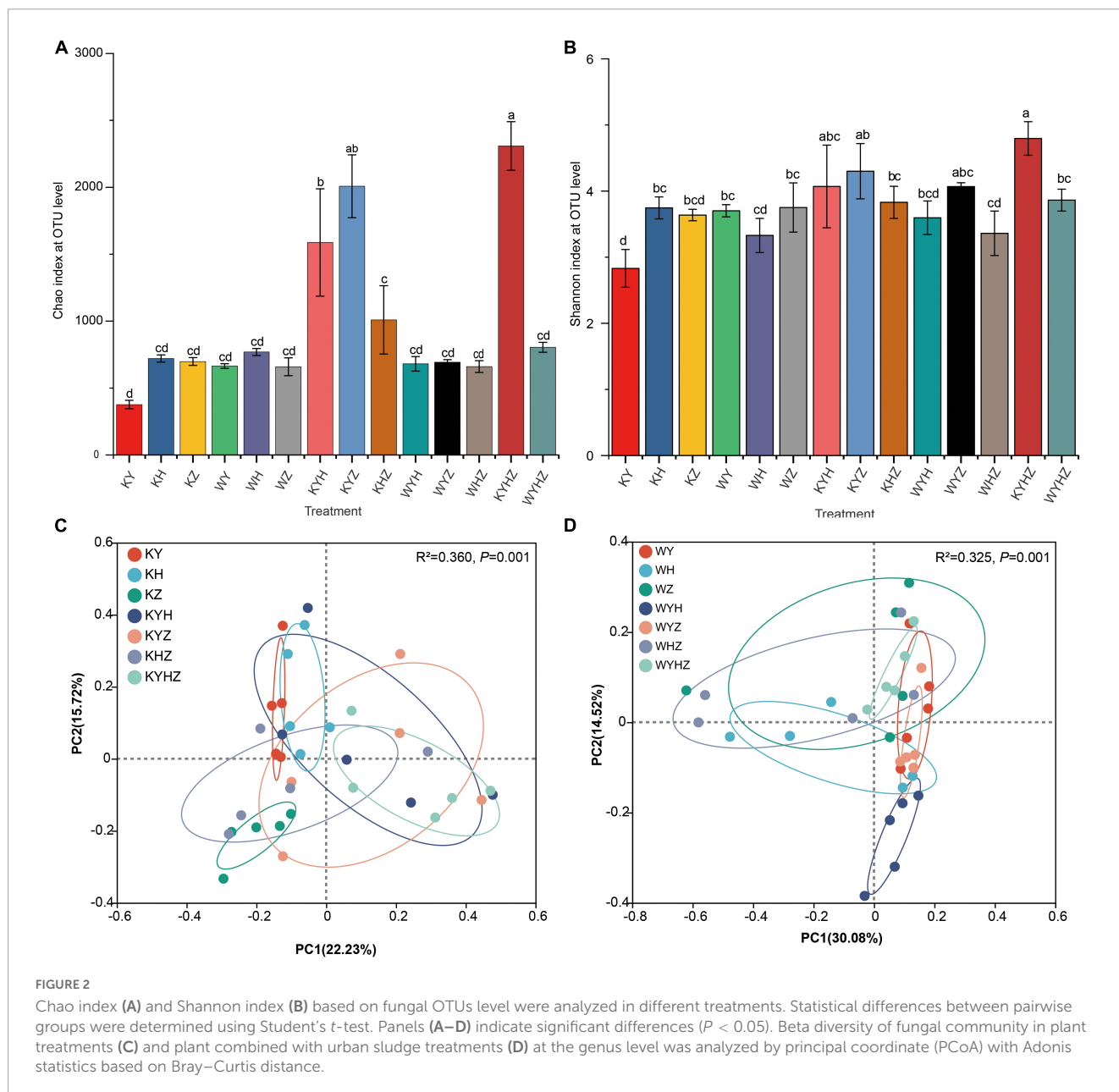
3.1. Soil properties and plant biomass

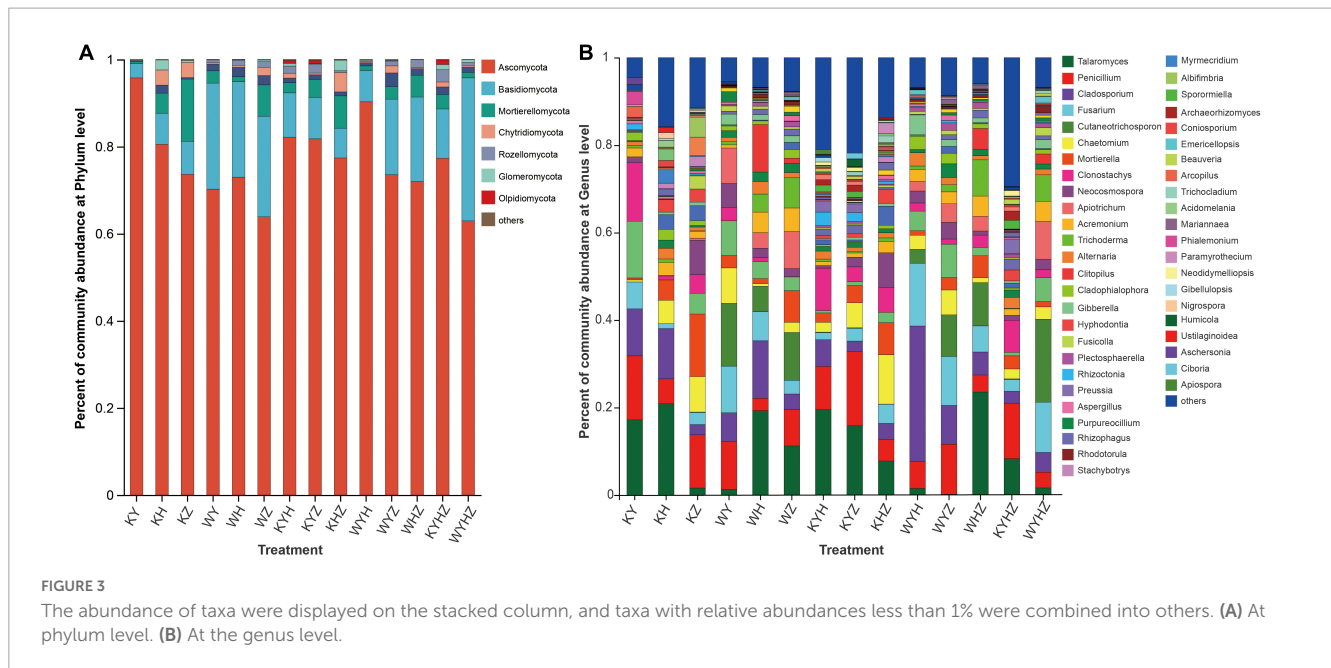
Soil properties were analyzed and compared in 14 treatments soil (Table 1). The content of AN and AP in plant treatments had

no significant difference compared with untreated quartz tailings soil. Notably, AN and AP of WH, WZ, WHZ, WYHZ, and AN of WY were significantly higher than in untreated quartz tailings soil ($r = 0.46, P < 0.05$). The content of AN and AP were increased with the addition of urban sludge. pH value of quartz tailings soil was increased after all treatments. In addition, the content of Fe_2O_3 in KH, KZ, KYH, KYZ, KHZ, KYHZ, WH, WZ, and WYHZ was significantly higher than that in untreated soil ($P < 0.05$), while the content of SiO_2 had no significant difference between treated and untreated tailings soil. The biomass of three plant species in plant combined with urban sludge treatments were higher than that in plant treatments (Supplementary Table 1) ($P < 0.05$). The highest biomass of Y, H and Z were in WYH, WYHZ, and WYZ, respectively. The total biomass of plant combined with urban sludge treatments was higher than in plant treatments ($P < 0.05$).

3.2. Effects of different treatments on fungal diversity and community composition

The soil fungal alpha diversity was analyzed based on Chao and Shannon index (Figures 2A, B). The fungal alpha diversity in KY was the lowest compared with KH and KZ ($P < 0.05$). Interestingly, the fungal alpha diversity significantly increased under Y combined with other plant species in plant treatments. However, the addition of urban sludge significantly decreased the fungal alpha diversity in the plant species combination treatments ($P < 0.05$). Notably, fungal alpha diversity in WYHZ and WYZ were higher than other treatments, respectively. Soil fungal community composition of KY, KH, KZ, and KYHZ were separated from each other while that of KYH, KYZ, KHZ, and KYHZ cannot be separated with each other at the genus level (Figure 2C). The fungal community





composition in WYH was different with other plant combined with urban sludge treatments (Figure 2D). Moreover, the plant treatments were separated from plant combined with urban sludge treatments.

The fungal community composition at the phylum and genus level were analyzed, Ascomycota, Basidiomycota, and Mortierellomycota were dominant phyla (Figure 3A). The addition of urban sludge decreased the abundance of Ascomycota and Mortierellomycota except for WYH, while increased the abundance of Basidiomycota. Additionally, the fungal community composition at the genus level among all treatments were significant difference (Figure 3B and Supplementary Figure 1). *Talaromyces*, *Penicillium*, *Clonostachys*, *Cladosporium*, *Mortierella*, *Chaetomium*, *Fusarium*, *Neocosmospora*, *Hyphodontia*, and *Acremonium* were dominant genera in plant treatments (Supplementary Figure 2A). However, the dominant genera in plant combined with urban sludge treatments were different with the plant treatments (Supplementary Figure 2B). The biomarkers in KYHZ and KZ were different with other plant treatments based on LEfSe analysis (LDA > 4) (Figures 4A, B). *Mortierella*, *Fusicolla* and *Stachybotrys* in KZ were dominant. *Cladosporium*, *Fusarium*, *Gibberella*, *Cladophialophora*, and *Alternaria* in WYH were dominant genera. Notably, the abundance of *Cutaneotrichosporon* and *Apiotrichum* were significantly higher in WYHZ than in other treatments.

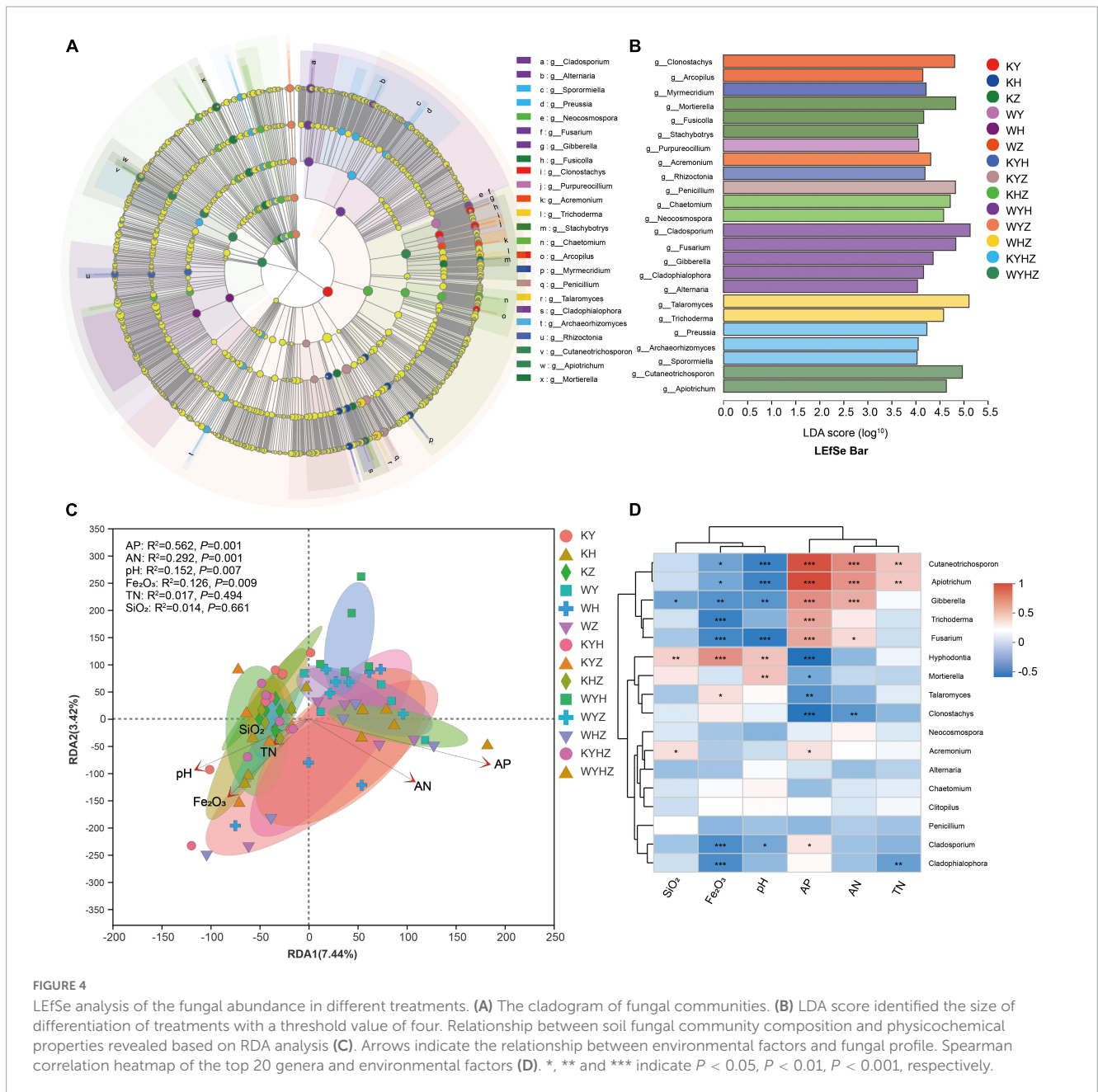
3.3. Effects of soil properties on fungal community composition

We analyzed the relationship between fungal community composition and soil properties using RDA analysis (Figure 4C). The results demonstrated that AP, AN, pH, and Fe₂O₃ significantly associated with the soil fungal community composition ($P < 0.01$) (Figure 4D). AP significantly affected on *Cutaneotrichosporon*, *Apiotrichum*, *Gibberella*, *Trichoderma*, *Fusarium*, *Acremonium*,

Cladosporium, *Hyphodontia*, *Mortierella*, *Talaromyces* and *Clonostachys*, and had negative correlation with the latter four genera. AN had significant positive correlation with *Cutaneotrichosporon*, *Apiotrichum*, *Gibberella*, and *Fusarium*, but negative correlation with *Clonostachys*. pH had significant positive correlation with *Hyphodontia* and *Mortierella*, and negative correlation with *Cutaneotrichosporon*, *Apiotrichum*, *Gibberella*, *Fusarium*, and *Cladosporium*. Moreover, Fe₂O₃ had a positive correlation with *Cutaneotrichosporon*, *Apiotrichum*, *Gibberella*, *Trichoderma*, *Fusarium*, *Cladosporium*, and *Cladophialophoron*.

3.4. Fungal network analysis

The results of the fungal network analysis showed that 3,574 and 277 edges were analyzed in the PT and PUT network, respectively (Table 2 and Figure 5A). PT network had higher total nodes, total links, connectedness, lower average path distance and modularity than PUT network. PT network was more complicated than PUT network. Furthermore, *Strelitziana* was network hub species, and *Didymella*, *Trichophaeopsis* were module hub species in PT network (Figure 5, Supplementary Figure 5, and Supplementary Table 2). *Fusarium* was network hub species, *Alternaria*, *Fusarium*, *Cerrena* were module hub species in PUT network. MEblue and MEBrown in PT network were negatively correlated with AP and AN, and positively correlated with pH (Supplementary Figure 3A). MEblue, MEBrown, MEBrown, MEgreen in PUT network were positively while MEyellow negatively correlated with AP and TN (Supplementary Figure 3B). MEblue was significantly correlated with MEBrown and MEyellow in PT network (Supplementary Figure 3C). MEBrown was significantly correlated with MEpurple, MEyellow, MEBlack, and MEblue in PUT network. MEblue, MEpink, and MEMagenta in PUT network were positively while MEBlack and MEBrown negatively correlated with pH (Supplementary Figure 3D).



MEblack and MEpurple in PUT network were positively correlated with AP, while MEpink was negatively correlated with AP ($P < 0.05$) (Supplementary Figure 3D). Key species in all modules were shown based on Module-EigenGene Analysis (Supplementary Figure 4).

4. Discussion

4.1. Effects of plant treatments on fungal diversity and community composition

In our study, soil fungal diversity in plant treatments significantly increased with plant species richness. The result supported our first hypothesis and consistent with previous

research (Gil-Martínez et al., 2021). The higher richness of the plant species may increase resource quantity of soil, and then increase fungal available niches (Waldrop et al., 2006; Cline et al., 2018). Moreover, soil fungal community composition in plant treatments was strongly driven by plant species (Figure 2). Plant can recruit their own fungal community as decomposer, mutualists, and inhibiting pathogens (Schlechter et al., 2019; Vives-Peris et al., 2020; Santoyo, 2021). Ascomycota that was the dominant phyla in all treatments can degrade cellulose and more complex carbohydrates and adapt to nutrient-poor and dry habitats (Lin et al., 2019; Shen et al., 2022). For example, *Preussia* and Archaeorhizomyces which were dominated in our samples. *Preussia* can produced glucosidases, phosphatases, cellulases, and IAA to promote plant development under stressful environmental conditions (Kandar et al., 2018, Toppo et al., 2022). Archaeorhizomyces can enhance

TABLE 2 Topological characteristics of the basic parameters in PT and PUT fungal network.

Fungal network	Node number	Edge number	Connectedness	Modularity	Average path distance
PT	295	3574	0.78	0.17	2.51
PUT	131	277	0.13	0.81	2.61

PT, plant treatments; PUT, plant combined with urban sludge treatments.

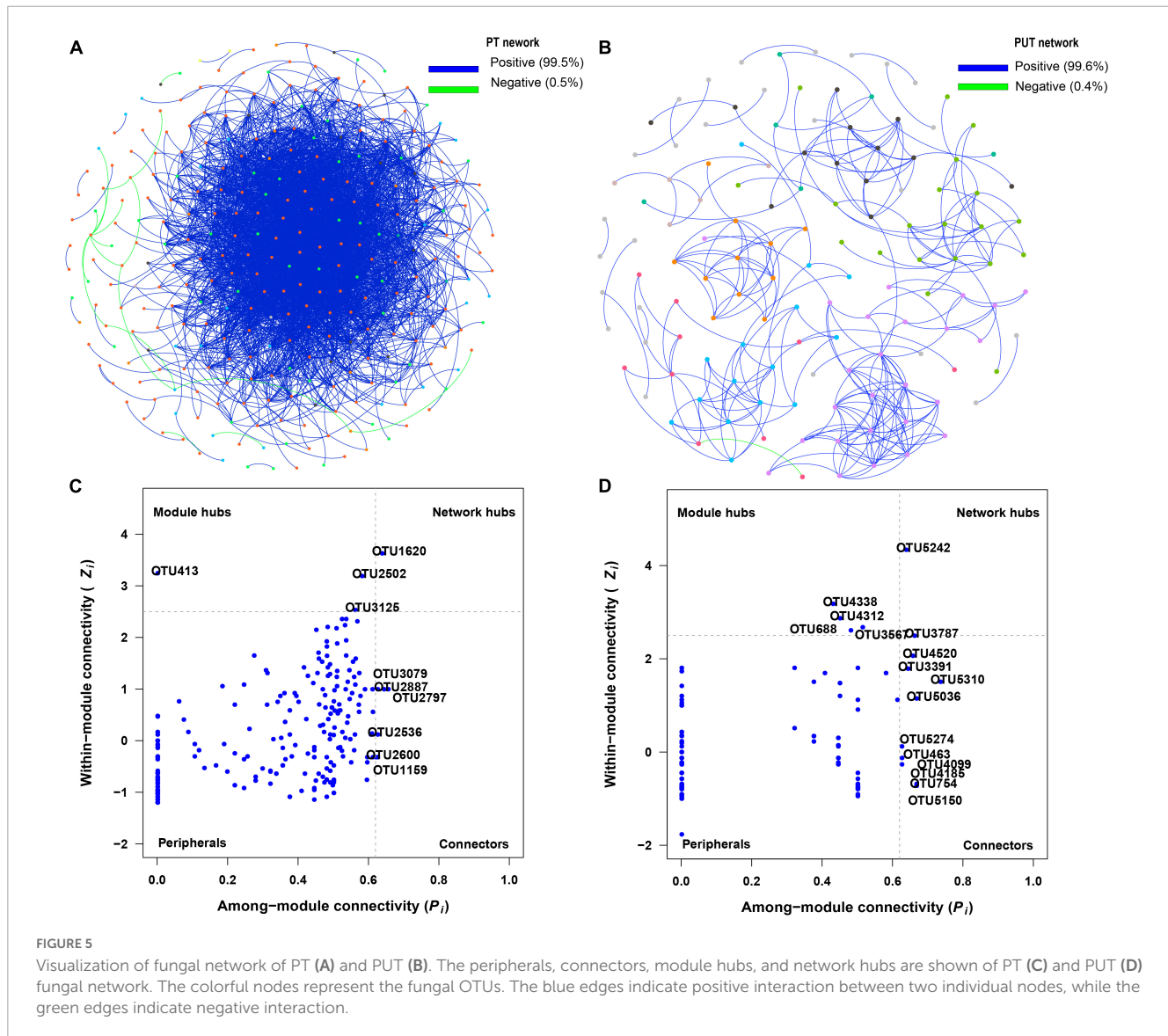


FIGURE 5 Visualization of fungal network of PT (A) and PUT (B). The peripherals, connectors, module hubs, and network hubs are shown of PT (C) and PUT (D) fungal network. The colorful nodes represent the fungal OTUs. The blue edges indicate positive interaction between two individual nodes, while the green edges indicate negative interaction.

plant bioactive components, improve stress resistance, and alleviate the occurrence of diseases (Zhang et al., 2020). Furthermore, *Mortierella*, *Fusicolla*, and *Stachybotrys* play an essential role in reducing the effects of environmental stresses on plant growth, preventing disease and even promoting bioremediation (Zhang et al., 2016; Li et al., 2021). *Cutaneotrichosporon* and *Apiotrichum* belonging to Basidiomycota were potential antagonist of soil-borne plant pathogens and plant growth promoter (Ortega et al., 2020; Buratti et al., 2022). Furthermore, *Apiotrichum* may degrade heavy metal pollution, and promote plant growth (Yalçın et al., 2018; Kumla et al., 2020; Xie et al., 2021). The results indicated that plant indirectly or directly promoted the fungal diversity and adopt the stressful environment of tailing soil through recruiting

potential beneficial fungal community (Wilkinson et al., 2019; Mattoo and Non-zom, 2021).

4.2. Variation and drivers of fungal community under plant combined with urban sludge treatments

In this study, soil properties, such as AN, AP, pH, TN, and Fe_2O_3 significantly affect soil fungal community composition. Furthermore, the addition of urban sludge significantly increased the content of AN and AP while decreased the fungal diversity

compared with plant treatments (Figure 2). The results suggested that AN and AP are the essential drivers in explaining the changes of fungal diversity, especially for acidic soil. Soil AN and TN mainly came from urban sludge and N-fixation of Legumes in 14 treatments. Our results showed that *Cutaneotrichosporon* and *Apiotrichum* were significantly associated with AN and TN. AN enrichment decreased soil fungal diversity due to toxic of more protons or weaken the linkage between soil carbon and fungal diversity (Yang et al., 2022c). Moreover, AN enrichment might aggravate the water limitation by promoting plant growth that were proved by the increase of plant biomass according our results. The water limitation may decrease some fungal survive (Angel et al., 2010). Moreover, AP positively and significantly associated with *Cutaneotrichosporon*, *Apiotrichum*, *Trichodema*, and *Gibberella* that were dominated in treatments. The results suggested that AP is important driver for fungal community composition (Bulgarelli et al., 2022).

Soil pH directly or indirectly effects fungal community composition in many ecosystems (Shen et al., 2020; Queiroz et al., 2021). Tailings soil with low pH is toxic for plant or soil microbial community. Our data showed that pH was increased after plant or plant combined with urban sludge treatments. The results suggested that pH was the crucial predictor for fungal community composition. After all, less fungal community can survive in lower pH soil. Thus, plant or organic matters in urban sludge mitigated soil acidification, and led to change of fungal community. Moreover, pH and Fe₂O₃ negatively and significantly associated with *Fusarium* indicated that can inhibit pathogen. Notably, the content of Fe₂O₃ increased after plant or plant combined with urban sludge treatments and then enhance plant biomass (Feng Y. et al., 2022). This may be related to that the increase of pH is conducive to the dissociation of iron oxides (Penn and Camberato, 2019). Notably, some fungi, such as *Talaromyces*, *Penicillium*, and *Cladosporium* can produce siderophores and then increase the content of Fe₂O₃ (Crowley, 2006; Pourhassan et al., 2014; Sahu and Prakash, 2021). Together, our findings on the changes of fungal community and soil properties after treatments supported that plant or plant combined with urban sludge effectively improve environment of tailings soil.

4.3. Fungal network under plant and plant combined with urban sludge treatments

In this study, the PT network had more node, edge numbers and complexity than the PUT network (Figure 5 and Table 2). More network connectors make the links stronger and make the network more stable. *Strelitziana* was network hub species, and *Didymella*, *Trichophaeopsis* were module hub species in PT network. The hub species were saprotrophs and mycorrhizal fungi that enhance plant nutrient uptake and development. AN was important driver of modular structure in PT fungal network. This suggests that fungal community tends to cooperate with each other to resist stress. Fungi-fungi feedback under low soil nutrition increases the stability and complexity of networks (Yang et al., 2022a). Furthermore, the addition of urban sludge increased modularity of network. In general, high modularity in the network was more beneficial to increase asynchronism, thus reducing the

influence of species loss on the overall network (Yang et al., 2022b). Moreover, the addition of urban sludge changed the soil properties, and AP, TN, pH, AN, and Fe₂O₃ significantly drive modular structure in PUT fungal network. Notably, *Fusarium* that are usually recognized as plant pathogens was hub species in PUT network. *Fusarium* has strong competitive ability by mycotoxin production (Karlsson et al., 2021). This suggests that pathogens may suppress other fungal groups and led to the decrease of stability and complexity of fungal network in PUT network.

5. Conclusion

In this study, we analyzed the fungal diversity and community composition under plant and plant combined with urban sludge treatments to remediate quartz tailings soil. The results suggest that plant diversity determine soil fungal diversity. Plant or plant combined with urban sludge significantly moderate acidification condition and increase nutrients of tailings soil. Plant can recruit beneficial fungi to improve their development and assist resist stress. Effectively remediate tailings soil due to the improvement of biological-chemical-physical properties. Fungal network is more stable and complex in plant treatments while fungal network is more fragile with the addition of urban sludge. AN and AP enrichment in urban sludge may be important drivers for fungal network. Future work should address the functional characters of fungal community when tailings soil is remediated by plant or urban sludge. Isolation and application of fungal functional groups are vital for remediation of tailings soil.

Data availability statement

The datasets presented in this study can be found in online repositories. The names of the repository/repositories and accession number(s) can be found in this article/[Supplementary material](#).

Author contributions

XL conceived and designed the experiments. FD analyzed the data and wrote the manuscript. YZ, XZ, and CZ analyzed the data. TS, YT, and YW conducted the pot experiments and measured soil parameters and plant biomass. All authors contributed to the article and approved the submitted version.

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Conflict of interest

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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Supplementary material

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fmicb.2023.1160960/full#supplementary-material>

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