

Effects comparison of different mulching methods on soil in pitaya orchards**

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Abstract. The aim of this study is to evaluate the effects of different mulching treatments, and to serve as a theoretical foundation for mulching practices in pitaya orchards of Hainan, China. In this study, the physicochemical properties, enzyme activity and also the structure and abundance of the soil microbial community were examined after a year's treatment. The results show that black fabric mulch and nature grass mulch significantly improved the soil water content, whereas its pH value was significantly reduced by purslane (*Portulaca oleracea* L.) mulch and black fabric mulch as compared to the control. Nature grass mulch increased soil organic carbon to a greater extent than control, coconut chaff mulch and also nature grass mulch significantly increased the content of alkaline hydrolysis nitrogen, nature grass mulch and black fabric mulch increased the content of available potassium more significantly than purslane (*Portulaca oleracea* L.) mulch and control. Treatments of nature grass mulch and purslane (*Portulaca oleracea* L.) mulch increased the activities of soil enzymes in the soil. The abundance of bacteria was highest in nature grass mulch and purslane (*Portulaca oleracea* L.) mulch, and the four mulch treatments helped to increase the richness of soil fungi more than control. The results show that all these types of mulch improved the characteristics of the soil compared with control, and nature grass mulch provided the best edaphic environment for pitaya orchards.

Keywords: mulching treatment, pitaya orchards, microbial community diversity, physicochemical characteristics of soil, Hainan province

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INTRODUCTION

Hainan province is one of the major producers of tropical fruits in China, and the pitaya industry is a pillar of the local economy. To mitigate the effects of extreme temperature, damaging sunlight, and the soil erosion during the growth of pitaya, an increasing number of local growers are using mulch in local orchards.

The types of mulch used in this area can be broadly classified into three groups: plastic film, organic material, and living mulch. Different mulching practices have varying effects on the physicochemical and biological properties of the soil. A large body of research has shown that mulching can increase the moisture content of shallow soil by reducing evaporation and regulating its temperature (Adamaviciene *et al.*, 2012; Ma *et al.*, 2018). Mulching treatments are also beneficial for controlling weeds, enhancing the content of soil nutrients, improving soil enzyme activities, and improving the microecological environment (Xun *et al.*, 2015).

While the conditions of the soil change according to the different mulching practices used, Chen *et al.* (2014) have claimed that although organic mulch is not mixed into soil, decomposed organic materials can be gradually absorbed into it. A relatively independent microecological system has been constructed by using film mulching which is more effective than organic mulching in terms of inhibiting evaporation and preserving heat in the soil (Zhang *et al.*, 2010). A study by Fang *et al.* (2011) showed that fresh biomass can

significantly improve the availability of nitrogen in the soil, and the supply of soil nutrients can be affected by the qualities of the mulching material. Living mulch can improve soil moisture by reducing evaporation (Unger and Vigil 1998), and can also improve the soil organic carbon (SOC) content (Xun *et al.*, 2015) by enhancing microbial activity (Yao *et al.*, 2005), which can help to decompose litter or mowed plants (Sanchez *et al.*, 2003). The particular effects of mulching treatments on microbial communities in the soil have also been researched (Dong *et al.*, 2017). Recent studies have shown that microorganisms in the soil can be directly influenced by mulching treatments or changes in the soil conditions (Chao *et al.*, 2016; Chen *et al.*, 2014).

The effects of mulching treatments vary significantly in different environments and climates, comparative studies have determined that due to its higher temperature and the effects of a higher precipitation rate, and also, a lower cation exchange capacity (CEC), the turnover of soil C in tropical soils is twice as rapid as it is in temperate soils (Six *et al.*, 2002). Also, mulching treatments can accelerate the loss of organic matter from the soil, thus further degrading it (Bot *et al.*, 2003). However, at present, there is a lack of research concerning orchard mulching with a focus on tropical regions, this is particularly the case in China. In our previous study, we compared two nearby pitaya orchards in Hainan province, the results show that black fabric cloth can lead to better fertilizer utilization efficiency, also, the abundance and diversity of fungi was higher when black fabric cloth was applied as compared with living mulch but the opposite occurred in the case of bacteria species, they were more abundant with the application of living mulch (Luo *et al.*, 2019). The objective of this study is to examine and compare the effects of four types of mulch: black fabric mulch (BFM), coconut chaff mulch (CCM), nature grass mulch (NGM), and purslane (*Portulaca oleracea* L.) mulch (PO) on the physical and chemical characteristics, enzyme activities, and the diversity and abundance of the microbial

community of the soil. The aim of this research is to provide a scientific basis for choosing suitable types of mulch in Hainan province.

MATERIALS AND METHODS

Field studies were carried out at the Wanzhong experimental orchard base in the town of Jianfeng (108°46' E, 18°40' N) in Hainan in southern China, which has a tropical ocean monsoon climate. The average annual air temperature was 25.5°C and the mean annual precipitation was 1347.5 mm, with more than 90% of it occurring from May to October. The average annual duration of sunshine was 2534 h. The soil at the experiment site was savanna soil. The pitaya trees in this orchard were planted in 2012 without mulching treatment and periodically weeded through the application of a herbicide. The pitaya cultivar chosen was Dahong, and the trees were planted with a spacing of 1.5 × 1.8 m.

In July 2017, five treatments (Table 1) were set up and sustained as follows: 1) control (CK), 2) black fabric mulch (BFM), 3) coconut chaff mulch (CCM), 4) nature grass mulch (NGM), 5) purslane (*Portulaca oleracea* L.) mulch (PO). A random block design of the five treatments was arranged, wherein each treatment was applied to three randomly arranged plots with each occupying an area of 100 m² (10 × 10 m) and these plots were separated by 1.5 m buffer strips.

Previous studies have shown that the properties of soil may be determined in order to compare the effects of different mulching treatments under the same conditions over the course of a year (Qu *et al.*, 2019). A one-year study period was also applied in this experiment, the experimental period commenced in July 2017 and ended in August 2018. During this period, 5:3:2 NPK was applied each month to achieve plant growth at a rate of 200 kg N ha⁻¹, and the watering practices and other management measures were consistent with those used by local farmers (maintaining the soil moisture content within 50-80%).

Table 1. Experimental design of different mulching management

| Mulching management | Treatment method |
|---------------------|--|
| CK | Herbicide was applied to control weeds without mulching (glyphosate was applied at the recommended rate of 1125 g ha ⁻¹ in July 2017, it was subsequently sprayed on new weeds at a small-scale every half month to a month). |
| BFM | Only black fabric mulch was applied. |
| CCM | 1.5 cm layer of coconut chaff mulch was applied. |
| NGM | Self-seeding weeds (mainly graminaceous) were grown freely around the plot, they were 95% carpeted, and were mowed once a month. The plants were spread out under trees after each instance of mowing. |
| PO | Purslane (<i>Portulaca oleracea</i> L.) was planted around the plots which were 80% carpeted with grass one year later. |

The soils were sampled under the five mulching treatments (CK, BFM, CCM, NGM and PO) in August, 2018. For each plot, five samples were randomly collected from the topsoil (0-15 cm) and inter-row areas which avoided the issue of border interferences, these samples were mixed to form one sample. The samples were placed in sterilized plastic bags. Each one was passed through a 2 mm sieve to remove crop residues, pebbles, and other foreign materials, and was divided into two subsamples. The first set of subsamples were air-dried and immediately stored at 4°C for chemical and enzyme analysis. The second set of subsamples were stored at -80°C for DNA extraction. Five samples from each plot were collected to determine the bulk density and moisture content of the soil.

The pH of the soil was measured with a glass electrode using a soil to water ratio of 2.5:1. The bulk density of the soil was determined as the mass of oven-dried soil, and fresh soil was dried in an oven at 105°C for 24 h to determine the moisture content.

Alkaline hydrolysis nitrogen (AN), and the available phosphorus (AP) and potassium (AK) were measured using alkaline hydrolysis diffusion, the Bray II method, and a flow auto-analyser, respectively (Zhang and Gong, 2012). The SOC was measured using the K_2CrO_7 - H_2SO_4 oxidation method (Zhang and Gong, 2012). The activities of enzymes in the soil were measured following the procedure proposed by Wu *et al.* (2013), urease activity was determined using the indophenols blue method, catalase activity was determined using UV spectrophotometry, acid phosphatase activity was measured using a method described by Tabatabai and Bremner (1969), and invertase activity was measured by titration, Guan *et al.* (1986). All of the above parameters were determined with 3 duplicates.

Soil DNA was extracted from a 0.5 g sample of composite fresh soil using PowerSoil™ DNA Isolation Kits (Qiagen) which were used according to the manufacturer's protocol, and the quantity and quality of the DNA were determined using a NanoDrop™ 2000 spectrophotometer (Thermo Scientific, USA).

A quantitative PCR (qPCR) system was used to determine the copy numbers of the 16S and 18S genes in the samples. The forward primer 341F (5'-CCTACGGGAGGCAGCAG-3') and reverse primer 534R (5'-ATTACCGCGTCTTGG-3') were used to target the 16S rRNA (Vasileiadis *et al.*, 2012), and the forward primer ITS1-f (5'-TCCGTAGGTGAACCTGCGC-3') and reverse primer 5.8S (5'-TCCTCCGCTTATTGATATGC-3') were used to target the ITS genes (Ciric *et al.*, 2010).

An ABI 7000 (PE Applied Biosystems) real-time PCR machine was used to conduct the qPCR test. Each reaction was performed in a 20 µL mixture containing 0.3 µL of each primer (0.4 mM), 12.5 µL of SYBR Premix Ex Taq™ (Takara, Dalian, China), 0.3 µL of ROX dye, 1 µL of template DNA, and 5.6 µL of double-distilled water. Each reaction was duplicated and contained negative sam-

ples (free from DNA) in order to monitor the sample for potential contamination. The thermal profile was 95°C for 5 min followed by 35 cycles at 95°C for 15 s, 55°C for 30 s, 72°C for 1.5 min, and a plate-reading step at 80.5°C for 30 s. In order to calculate amplification specificity, a melting curve analysis was performed to confirm product specificity. The gene fragments of bacterial (V3-V4 region of the 16S rRNA genes) and fungal (ITS2 region of the 18S rRNA genes) origin were PCR-amplified and cloned into the pGEM-T Easy Vector. The clones that had the appropriate gene inserts were chosen as the standards for qPCR. Ten-fold serial dilutions of a known copy number of plasmid DNA with the appropriate gene inserts were generated to produce a standard curve, respectively. The bacterial and fungal copy number were expressed in terms of per gram of soil (fresh weight), the relative values were log-transformed and the R^2 value were >99%.

The V3-V4 region of the 16S rRNA genes was amplified by PCR according to the bacterial primers 341F (5'-CCTACGGGNGGCWGCAG-3') and 805R (5'-GACTACHVGGGTATCTAATCC-3'), and the ITS2 region of the 18S rRNA genes was amplified using the fungal primer BITS (5'-ACCTGCGGARGGATCA-3') and reverse primer B58S3 (5'-GAGATCCRTTGYTRAAAGTT-3') (Bokulich and Mills, 2013).

A 25 µL PCR reaction featuring 1 µL of the template DNA, 15 µL of 2 × Taq PCR MasterMix (Aidlab), 0.5 µL of forward and reverse primers (5 µM of each primer), and 8 µL of double-distilled water was performed for every sample. The PCR conditions were as follows: 94°C for 45 s, 30 cycles of 95°C for 30 s, 54°C for 30 s, and 72°C for 30 s, followed by 5 min at 72°C. The products of PCR were extracted and purified after amplification. The amplicons were then quantified and pooled at equal concentrations, and diluted to create one sample. Sequencing was performed using the Illumina HiSeq PE250 platform at the Personal Biotechnology Co., Ltd. Shanghai, China.

The bacterial and fungal sequencing data, with the adaptor and primer sequences removed, and a unique barcode assigned to each sample were analysed by the Quantitative Insights into Microbial Ecology (QIIME) software package (version 1.7.0). The sequences covered the V3-V4 region of the bacterial 16S rRNA genes, and the ITS2 region of the fungal 18S rRNA genes were clustered into operational taxonomic units (OTUs) at a confidence threshold of 97% by using UPARSE software (version 7.1, <http://drive5.com/uparse/>) (Edgar 2013). The OTUs after this were used to annotate taxonomic information. Finally, the sequences in this study were sent to the Sequence Read Archive (SRA) database of the NCBI under accession number PRJNA574235 for the bacteria and fungi.

Alpha diversity (Chao1, ACE, Shannon, and species' diversity) was calculated using QIIME (version 1.7.0). The taxonomies of the 16S rRNA gene sequence and 18S rRNA sequence were analysed using the Ribosomal Database

Project (RDP) classifier for each sample. The IBM SPSS 19.0 software and Excel 2010 were used to perform a statistical analysis. Significant differences in the characteristics of the soil, the activities of the enzymes in it, and OTU abundance (at $p < 0.05$) were among the treatments that were assessed using an ANOVA with Duncan's multiple range test. The correlation between the characteristics of the soil, the activities of the enzymes in it and the soil samples subjected to different treatments were determined by redundancy analysis (RDA) using CANOCO 5.1 software (Etten, 2005).

RESULTS

The influence of mulch application on the physical and chemical characteristics of the soil are shown in Table 2. The soil moisture content was the lowest for CK (7.93%) and the highest for BFM (9.84%). The bulk density of the soil was highest for BFM, followed by CK and CCM, and had its lowest value for PO. The pH of the soil for PO was significantly lower than it was for CK, CCM, and NGM ($p < 0.05$).

The results show that the content of soil nutrients were higher for NGM than for CK, and the available nutrients were higher for BFM and CCM than for CK. The different mulching modules improved the soil nutrient content to varying degrees. The SOC values of samples treated with NGM were higher than those for CK, and the differences between BFM, CCM, and PO were not significant ($p > 0.05$). The contents of soil AN in CCM and NGM were

significantly higher than it was in BFM ($p < 0.05$). The AP content was higher in NGM than in CCM and CK. AK content was higher in BFM and NGM than in PO and CK.

The soil catalase activities in NGM were higher than in PO, BFM and CK (Table 3). The acid phosphates and invertase activities of soil were higher in NGM than in the other treatments, to a significant extent. Acid phosphate activities were the lowest in CK and BFM as compared to other treatments. The soil urease activities were higher in NGM, PO and CCM than in CK and BFM ($p < 0.05$).

There were no significant differences in bacterial OTUs, while the number of fungal OTUs in PO was lower than in other treatments ($p < 0.05$) (Table 4). The coverage of these treatments ranged from 95.13 to 96.19% for bacteria, and from 99.75 to 99.84% for fungi.

Alpha diversity analysis was used to examine the abundance and diversity of the microbial communities in the soil (Table 4). For bacteria, the ACE index was higher for PO than for CK. There was no prominent difference in the Chao1 index between the various treatments. The Shannon index was significantly higher for all four mulch treatments than it was for CK ($p < 0.05$). The results show that the mulch treatments helped to increase the degree of diversity and richness of bacteria in the soil compared to CK. For fungi, the ACE index and the Chao1 index were lower for PO than for other treatments.

And the Shannon index was lower for PO than for BFM, CCM and NGM ($p < 0.05$).

Table 2. Physicochemical analysis of soil subjected to different mulch treatments

| Mulching management | SOC (g kg ⁻¹) | AN | AP | AK | Moisture content (%) | Bulk density (g cm ⁻³) | pH (H ₂ O) |
|---------------------|---------------------------|------------------------|--------------|---------------|----------------------|------------------------------------|-----------------------|
| | | (mg kg ⁻¹) | | | | | |
| CK | 17.43±1.25b | 33.97±1.29c | 13.52±0.46c | 161.33±2.62c | 7.93±0.85c | 1.36±0.02b | 5.94±0.05a |
| BFM | 18.08±1.11ab | 39.69±2.25b | 19.05±0.24ab | 206.33±5.51a | 9.84±0.36a | 1.49±0.03a | 5.47±0.20bc |
| CCM | 19.14±0.87ab | 51.07±3.40a | 17.46±0.68b | 190.33±9.29ab | 9.00±0.19ab | 1.37±0.03b | 5.90±0.16a |
| NGM | 20.62±1.12a | 48.29±1.35a | 20.11±0.87a | 202.33±10.79a | 9.15±0.20ab | 1.32±0.04bc | 5.81±0.02ab |
| PO | 18.09±1.48ab | 37.19±1.55c | 18.97±0.85ab | 176.00±3.61bc | 8.03±0.29bc | 1.27±0.01c | 5.25±0.12c |

CK – control, BFM – black fabric mulch, CCM – coconut chaff mulch, NGM – nature grass mulch, PO – purslane (*Portulaca oleracea* L.) mulch. Mean ± standard error.

Table 3. Catalase, acid phosphates, urease and invertase activities in samples subjected to different mulch treatments

| Mulching management | Catalase activity | Acid phosphate activity | Urease activity | Invertase activity |
|---------------------|----------------------|-------------------------|-----------------|--------------------|
| | (U g ⁻¹) | | | |
| CK | 1.05±0.06c | 0.87±0.08d | 10.37±0.64b | 37.13±0.62cd |
| BFM | 0.65±0.04c | 1.01±0.02d | 9.43±0.61b | 35.34±4.44d |
| CCM | 2.08±0.13ab | 1.55±0.04c | 12.83±0.87a | 42.54±0.39bc |
| NGM | 2.71±0.15a | 2.61±0.02a | 13.70±0.46a | 57.25±0.35a |
| PO | 1.36±0.59bc | 2.27±0.14b | 13.63±0.72a | 47.93±1.12b |

Mean ± standard error, values followed by different letters differ significantly (Duncan's test, $p < 0.05$). Other explanations as in Table 2.

Table 4. Alpha diversity measures of bacteria and fungi in samples subjected to different mulch treatments

| Mulching management | Coverage (%)* | OTUs | ACE | Chao1 | Shannon |
|---------------------|---------------|-----------|------------|-----------|-------------|
| Bacteria | | | | | |
| CK | 96.19±0.31 | 6058±169a | 8716±83b | 8559±100a | 10.42±0.02b |
| BFM | 95.15±0.76 | 6235±673a | 9281±775ab | 9065±827a | 10.70±0.06a |
| CCM | 95.31±0.74 | 6146±308a | 9107±199ab | 9019±249a | 10.65±0.02a |
| NGM | 95.63±0.24 | 6075±68a | 9044±66ab | 8879±81a | 10.66±0.05a |
| PO | 95.13±0.84 | 6842±648a | 10080±659a | 9846±673a | 10.74±0.06a |
| Fungi | | | | | |
| CK | 99.84± 0.01a | 2334±188a | 3304±207a | 3226±198a | 6.82±0.85ab |
| BFM | 99.76±0.00b | 1993±131a | 2726±242b | 2717±250a | 6.92±0.22a |
| CCM | 99.80±0.02ab | 2152±150a | 3019±194ab | 2941±171a | 7.19±0.03a |
| NGM | 99.81±0.01ab | 2247±115a | 3163±167ab | 3095±187a | 7.06±0.08a |
| PO | 99.75±0.06b | 1112±190b | 1470±225c | 1520±203b | 5.72±0.38b |

Mean ± standard error. Values followed by different letters differ significantly (Duncan's test, $p < 0.05$). *For library coverage of samples. Other explanations as in Table 2.

The relative abundance of the bacterial and fungal communities at the phylum level are shown in Fig. 1. For bacteria, the first 10 phyla are shown, the most dominant phyla were *Proteobacteria* and *Actinobacteria*, accounting for more than 45% in all five treatments. The relative abundance of *Proteobacteria* was 37.7, 36.6, 37, 38.2 and 31.7% for BFM, CCM, NGM, PO and CK. *Actinobacteria* were found to be at a level of 18.9% in CK, and were 17.6, 15.1, 14.5 and 11.7% in NGM, PO, CCM and BFM, respectively. The third most abundant phylum (10% relative abundance on average) was *Acidobacteria*. This phylum was the most abundant one in samples treated with CCM (11.1%) followed by CK (10.7%), BFM (9.9%), NGM (9.6%), and PO (8.6%). The next most abundant phyla were *Gemmatimonadetes*, *Planctomycetes*, and *Chloroflexi*, the relative abundance of each of these was greater than 5%.

For fungi, *Ascomycota* was the dominant phylum across all soil treatments. *Basidiomycota* and *Mortierellomycota* only accounted for a minor part of the total. The relative abundance of *Basidiomycota* ranged from 2.0 to 4.0%. The relative abundance of *Mortierellomycota* was 3.5% in BFM.

The bacterial and fungal abundance was quantified using qPCR, and the results show a significant difference between samples subjected to different mulch treatments. Bacterial abundance ranged from 0.48×10^6 to 1.82×10^6 copy numbers/g of fresh soil (Fig. 2a). The decreasing order of treatments in terms of the copy numbers of the 16S rRNA gene is NGM>PO>CCM>BFM>CK. The bacterial abundance in samples subjected to NGM and PO was 1.82×10^6 and 1.59×10^6 copy numbers/g fresh soil, respectively, it was significantly higher than those of other treatments ($p < 0.05$). For fungi, the average abundance ranged from 0.12×10^6 to 1.57×10^6 copy numbers/g of fresh soil for different mulch treatments (Fig. 2b). The sequence

of abundance of 18S rRNA from the highest to lowest was NGM>PO>CCM>BFM>CK. Samples subjected to NGM had the largest fungi population, it was 13.08 times higher than that of CK.

Redundancy analysis (RDA) was used to analyse the relationships between the various characteristics of the soil, and the bacteria and fungi in it with different treatments (Fig. 3). For bacteria, the first and second components explained 64.03 and 24.28% of the total variation in orders, respectively. The CK treatments were separated from all of the other treatments in case of bacterial data. The activities of catalase, invertase, urease, and phosphate demonstrated close correlations between the bacterial community. The bulk density of the soil and its water content were predominantly associated with CCM and BFM, and also, the pH value was positively correlated with CK (Fig. 3a).

For fungi, the first axis of the RDA explains 93.54% of the total variation. Urease, invertase, and phosphates were positively correlated with PO, and catalase was positively correlated with CCM. The bulk density of the soil, the moisture content, and AK showed close correlations with BFM. The three samples subjected to CK were separated from one another in the fungal data (Fig. 3b).

DISCUSSION

The four mulching treatments (BFM, CCM, NGM, and PO) had various influences on the physical and chemical properties of the soil. For example, the moisture and bulk density of the soil were highest in BFM, which is consistent with past work (Liu *et al.*, 2014a). Previous studies reported that black plastic film mulch is more efficient than organic mulch at reducing the rate of evaporation (Zhao *et al.*, 2019). This might be why the bulk density of soil was significantly higher for BFM than for other mulch treatments.

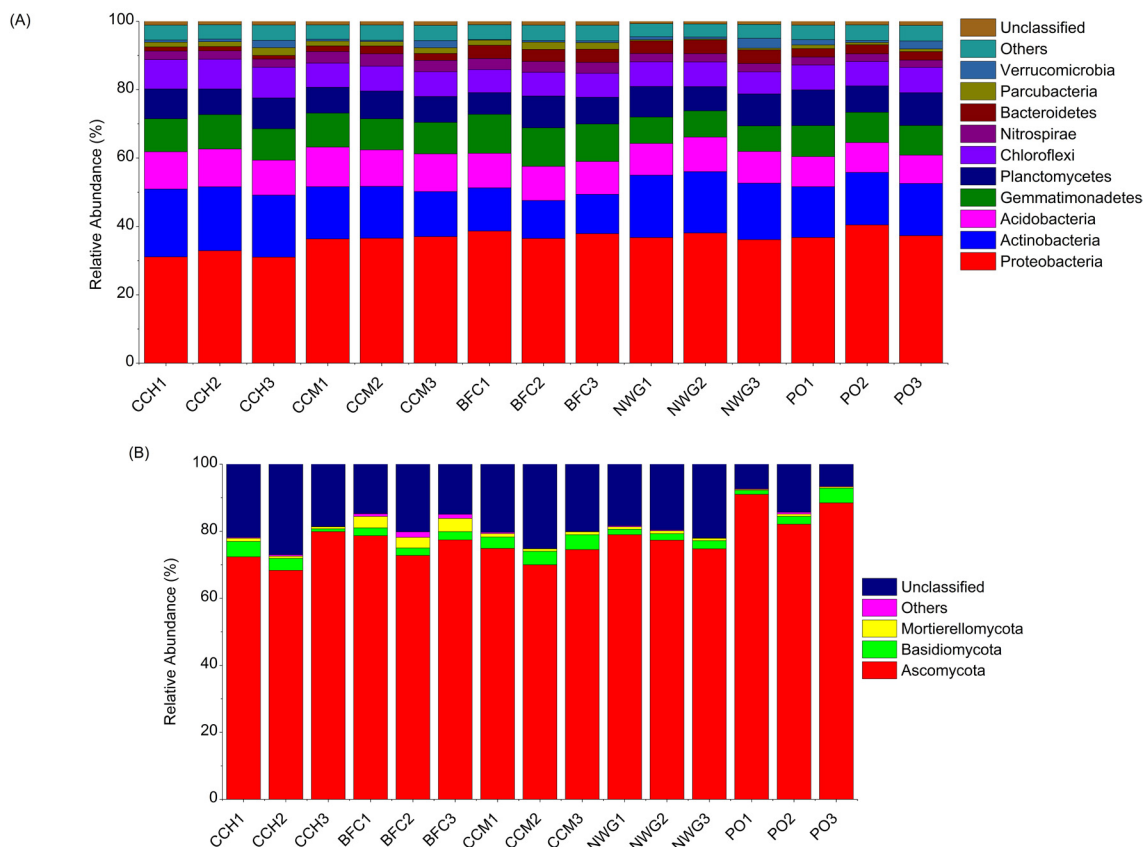


Fig. 1. Relative abundance of bacterial (A) and fungal (B) phyla in soil subjected to different mulch treatments. CK – control, BFM – black fabric mulch, CCM – coconut chaff mulch, NGM – nature grass mulch, PO – purslane (*Portulaca oleracea* L.) mulch.

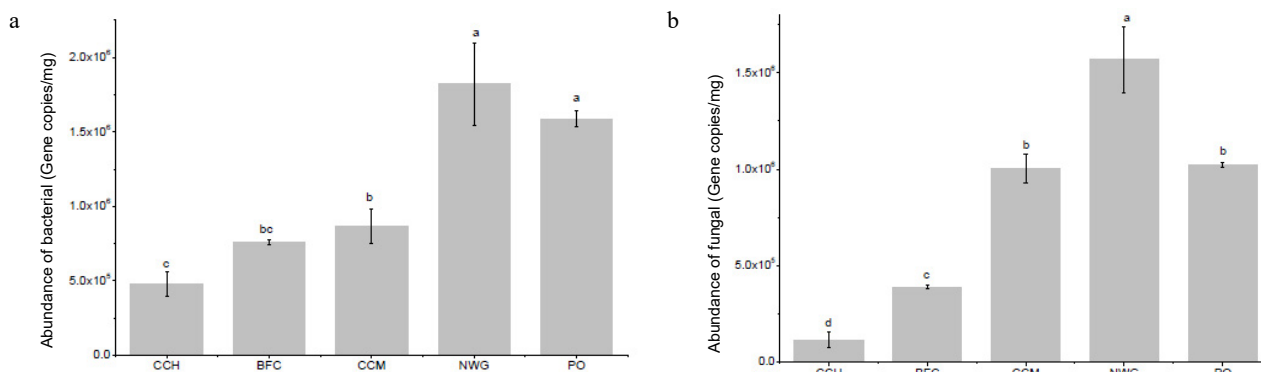


Fig. 2. Abundance of bacteria (a) and fungi (b) in the soils of the pitaya orchard under different mulching patterns. CK – control, BFM: black fabric mulch, CCM – coconut chaff mulch, NGM – nature grass mulch, PO – purslane (*Portulaca oleracea* L.) mulch. Values followed by different letters differ significantly (Duncan's test, $p < 0.05$). Bars represent standard errors.

The SOC content was lower for CK than for NGM. Other studies found that SOC is not significantly higher for BFM than for CK (Ma *et al.*, 2018; Luo *et al.*, 2019), which is consistent with the results of this study. Previous research has also shown that the long-term application of a film of mulch reduces the SOC content of soil (Xiaobo *et al.*, 2018), this is possibly due to increased mineralization (Xiao-Gang and Feng-Min, 2015). Cadavid *et al.* (1998) have reported that organic mulch can significantly benefit by increasing the SOC, but in our research, CCM did not have a significant effect on SOC, possibly because of the

short mulching period. Each type of mulch except for PO significantly improved the availability of soil nutrients. Qu *et al.* (2019) have also shown that different mulch treatments have similar effects concerning the improvement the available nutrients in soil. The contents of SOC and available nutrients in NGM were higher than those of CK. However, in the case of SOC, AN and AK there was no clear increase in PO compared with CK, which indicates that naturally occurring grass and purslane (*Portulaca oleracea* L.) yielded different results in terms of the contents of nutrients in the soil. Zhong *et al.* (2018) also reported that

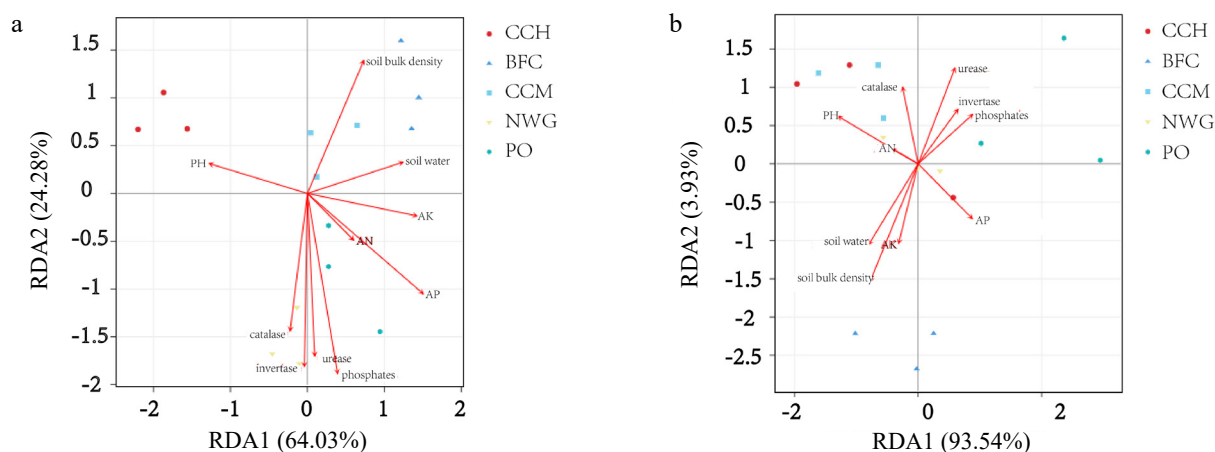


Fig. 3. Redundancy analysis (RDA) of the compositions of bacterial (a) and fungal (b) communities, and their environmental variables with different mulch treatments. CK – control, BFM – black fabric mulch, CCM – coconut chaff mulch, NGM – nature grass mulch, PO – purslane (*Portulaca oleracea* L.) mulch.

the effects of improving the soil nutrients vary between the living mulch species. A study concerning living mulch has shown that mulch plants can increase the content of nutrients in the soil by improving its microbial activity (Yao *et al.*, 2005) and producing root exudates (Hoagland *et al.*, 2008). However, in another way, mulch plants can reduce nutrients in soil by absorbing them (Xun *et al.*, 2015). In this study, the available nutrient contents were higher for NGM than for PO, which shows that grass types have an obvious influence on the availability of nutrients in the soil (Zhong *et al.*, 2018).

The activity of soil enzymes is the key factor influencing biochemical processes in the soil (Xiaobo *et al.*, 2018), and can make a contribution to maintaining or even increasing soil fertility (Burns *et al.*, 2013). For example, urease and acid phosphatase in the soil have been shown to be related to such nutrients as those involved in the mineralization of N, P, and K (Haroun, 2010). Other studies have shown that the activities of enzymes in the soil are enhanced in rooted soil by the plant-mediated activation of microorganisms (Cheng *et al.*, 2005). Different types of grass have different effects concerning the enhancement of the activity of soil enzymes. For example, studies have shown that leguminous mulch can improve soil enzyme activity more efficiently than Gramineae mulch (Adamaviciene *et al.*, 2012). Caravaca *et al.* (2005) found that the values of hydrolases were highest in *H. portulacoides* rhizosphere soil. In this research, NGM contributed more significantly to improving soil enzyme activity than PO, which also shows that the relationship between the species of grass applied to the soil and the enzyme activity was significant. The activities of enzymes in CCM was significantly higher than that in CK. Some scholars claim that mulching treatments can improve the moisture content of the soil by reducing the rate of evaporation (A'Bear *et al.*, 2014). There was no difference

in the enzyme activities of catalase, urease, and invertase between BFM and CK, the tropical heat probably had the same effect in both cases.

The related experiments show that mulching types affect both microbial abundance and composition (Wei *et al.*, 2018b), and microorganisms in the soil can enhance the cycling of nutrients and improve their storage prospects (Shibahara *et al.*, 1998). In our research, the diversity and richness of soil bacteria were higher in all mulch-treated samples, as found in previous studies (MeiyanWu *et al.*, 2009). Jin *et al.* (2009) have suggested that the microbial population is related to the influence of mulch on the physical and chemical characteristics of the soil. In the case of fungi, NGM had a higher richness than other treatments, Schirmel *et al.* (2018) demonstrated that the activities of soil microbes is significantly higher in organically mulched fields than in fields with plastic mulch. Pan *et al.* (2016) found that organic mulch helps to promote the microbial biomass of soil and increase the proportion of fungi. The improvement in the conditions of soil are important in terms of improving the abundance and diversity of the microorganisms within it (Mbah *et al.*, 2014). The improvement in certain characteristics of its physical-chemical structure, such as the organic matter content, nutrient availability, moisture status, and porosity in organic mulches, helps to improve the growth of microorganisms (Abdullah, 2014). Some research suggests that inefficient nutrient cycling in plastic mulch can reduce the level of soil microbial activity (Schirmel *et al.*, 2018). Numerous studies have shown that the litter properties of plant species are essential to the structure and activity of the soil microbial community (Bardgett, 2007). While PO produced the lowest value of fungal diversity, previous studies have shown that differences in bacterial and fungal diversity in soil may be explained through the variation in plant residues and the input of exudates between plant species (Legay *et al.*, 2014). Previous studies have also suggested that

Actinobacteria and *Acidobacteria* show a clear preference for growing in nutrient-poor soil (Zhang *et al.*, 2016). The phyla of these two bacteria were dominant in our research, which is consistent with the results of past studies (Chao *et al.*, 2016). Wang *et al.* (2019) concluded that a decrease in soil nutrient content can stimulate an increase in the number of branches of *Proteobacteria* and *Actinobacteria*. The *Actinobacteria*, which can decompose recalcitrant organic carbon (Dang *et al.*, 2017), are spore-forming bacteria that dominate in harsh and stressful soil conditions. It has been established that *Acidobacteria* tend to be more proficient with regard to soil C transformation and the metabolization of recalcitrant organic substrates (Li *et al.*, 2017), also, some subgroups of *Acidobacteria* have been observed to be abundant in soil with a higher SOC content (Liu *et al.*, 2014b). Other research has concluded that *Acidobacteria* are more abundant oligotrophic organisms in carbon-poor soils (Wang *et al.*, 2019; Noah *et al.*, 2007) also found that *Acidobacteria* were relatively less abundant in resource-rich plots than in no-litter plots, which may be explained by the copiotrophic/oligotrophic model of community effects on decomposition. It was found that the fungal diversity of soil was directly related to its properties (such as its physical-chemical characteristics) (Yang *et al.*, 2017). *Ascomycota* was the dominant phylum of all samples subjected to treatment in this research, and is also the dominant fungal community in the majority of ecosystems worldwide (Menkis *et al.*, 2015). Many taxa belonging to *Ascomycota* are associated with the decomposers of organic matter (Urbina *et al.*, 2016), and also, organic mulch might facilitate the growth of this phylum. The properties of the plants growing in the soil have an indirect effect on fungal diversity (Yang *et al.*, 2017), indeed several studies have suggested that fast-growing plant species that produce large amounts of litter and root exudates are more effective than slow-growing species in terms of enhancing fungal diversity (Kembel *et al.*, 2012). The relationship between the microorganism community in the soil and its physicochemical conditions remains unclear (Wei *et al.*, 2018a). Our study indicates shifting patterns in the structure and community of the microbial biomass under different mulch treatments.

CONCLUSIONS

1. The soil water content for black fabric mulch was significantly higher than it was for purslane (*Portulaca oleracea* L.) mulch and control, whereas the bulk density increased due to the application of black fabric mulch. purslane (*Portulaca oleracea* L.) mulch was favourable due to the decrease in soil bulk density.

2. Different mulching modules improved the soil nutrient content to varying degrees. Soil nutrients such as soil organic carbon, alkaline hydrolysis nitrogen, available potassium and available phosphorus were higher for nature grass mulch than for control.

3. Coconut chaff mulch, nature grass mulch and purslane (*Portulaca oleracea* L.) mulch were helped to improve the activities of urease and acid phosphate. The invertase activity was highest for nature grass mulch as compared to the other treatments.

4. Moreover, mulch treatments can help to improve the abundance of bacteria and fungi. The changes in the bacterial and fungal communities were correlated with the physicochemical properties of the soil, which might be affected by mulching practices.

5. A long-term assessment of the soil environment is required to obtain a clear understanding of different mulching practices.

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