Loss of Microbial Diversity exacerbates the Spread of Antibiotic Resistome in Soil

Abstract: Biodiversity loss is considered as a major threat because of its importance for ecosystem processes and services on which society depends. Natural ecosystems show variable resistance to invasion by alien species, and this resistance can be related to the microbial diversity in the system. For centuries, little has been known regarding the role of soil microbial diversity in the biological barrier against microbial invasion. The increasing prevalence of antibiotic resistant bacteria is one of the most serious threats to public health in the 21st century. Here we explore whether reductions in soil microbial diversity have consequences on the dissemination of antibiotic resistance. By using a dilution-to-extinction approach coupled with a high-capacity quantitative PCR arrays, we investigated the relationship between soil microbial diversity and the barrier effect against antibiotic resistance invasion. We observed a negative correlation between microbial diversity and the abundance of antibiotic resistance genes (ARGs), and this correlation was maintained even when accounting simultaneously for other drivers (incubation time, microbial abundance). Our results demonstrated that higher microbial diversity can act as a biological barrier against antibiotic resistance spread. Together with previous work, our results fill a critical gap in understanding the role of soil microbial diversity in ecosystem health

Introduction

It is well documented that biodiversity is the foundation of the maintenance of ecosystems, i.e. a large species pool is required to sustain a health ecosystem since they play a key role in multiple ecosystem functions and services simultaneously, including earth's biogeochemical cycling, primary production, litter decomposition as well as climate regulation [1-4]. The rapid and continued development of molecular biology and genomic techniques has unveiled immense microbial diversity in soil and ocean [5-8]. However, the roles of microbial diversity and associated traits in controlling ecosystem functioning remain unclear [9-11].

In recent years, a few studies have shown that anthropogenic activities (such as agricultural intensification, land use and nitrogen enrichment), and climate change may reduce microbial diversity, a response that will likely impact ecosystem functions [12-15]. There is an extensive debate about relationships between microbial diversity and ecosystem functioning [16, 17]. Currently there are three different views about the relationships. It has been proposed that soil microorganisms are key components that determine life supporting functions, but functional redundancy in soil microorganisms are prevalent, therefore an initial loss in microbial diversity is unlikely to substantially affect ecosystem functions[16, 18, 19]. Studies have also demonstrated that the relationships between microbial diversity and ecosystem functions are saturated [20], suggesting losses of a few species at high richness levels could have minimal consequences on the ecosystem functioning [21]. For instance, microbial decomposer communities often

39   exhibit high redundancy for a single function, such as microbial respiration and biomass

40   production [20, 22]. However, if the function of concern was more specific than general

41   functions, such as pesticide degradation are known to be limited to only a few

42   specialized functional groups[23], the relationship between biodiversity and ecosystem

43   functioning is more linear than saturating [24]. Moreover multifunctional redundancy was

44   generally lower, i.e. the degree of multiple functional dependence on diversity was

45   higher than single-functional redundancy [25]. Therefore, the buffering capacity of

46   ecosystem functions against biodiversity loss may be limited that a moderate loss of

47   diversity may substantially impair key specialized functions [23]. Additionally, Jung et al.,

48   observed that genes related to the nitrogen cycling was significantly reduced,

49   contradictory results have also been obtained that the efficiency of diesel

50   biodegradation was increased in the low-diversity community [26]. These results

51   indicated that the relationship between microbial diversity and ecological function

52   involves trade-offs among ecological processes, and should not be generalized as a

53   positive, neutral, or negative relationship [26]. Each point of view sound reasonable, and

54   this could be, at least partly, the reason why the consequences of decline in biodiversity

55   for ecosystem processes and functioning have long been of considerable interest[27-30].

56   Resistance to invasion by alien species represents a major life support function of

57   terrestrial ecosystems[31, 32]. The biological invasion, developed in the 1950s, which

58   indicated that ecosystems that contain a higher level of biodiversity are less vulnerable

59   to disturbances [33-35]. Whether this theory can be applied to microbiology level remains

60 largely unresolved. To answer this question, some pioneering studies have investigated

61 the relationship between invasibility and the microbial diversity of ecosystems, by

62 manipulating the microbial community [19, 30, 36-38]. For example, van Elsas et al.,

63 observed that a positive correlation between the inoculant survival rate of *Escherichia*

64 *coli* O157:H7 and the soil fumigation depths; additionally, by using a dilution–

65 reinoculation approach they obtained a similar result that a negative correlation between

66 the soil microbial diversity and survival of the invader [36, 37, 39]. An inverse relationship

67 was also found between the survival rate of the inoculant and the degree of microbial

68 diversity with respect to *Pseudomonas aeruginosa*, *Ralstonia solanacearum* and

69 *Listeria monocytogenes*[39-41]. These results shed lights on the complexity–invasiveness

70 relationship within microbial communities. However, with investigating limited species,

71 we still lack a sound evidences to achieve a universal phenomenon. To address this

72 question in an alternative way, we focus on the relationship between invasibility and

73 microbial diversity at gene-level.

74 Antibiotic resistance genes (ARGs) were chosen as the invaded genes. On one hand,

75 antibiotic-treatment failure is typically attributed to the "weapon-shield" role that ARGs

76 played in clinical settings [42]. On the other, the pace of development of novel antibiotics

77 is now alarmingly low [43]. Now the emergence and spread of antibiotic resistance has

78 become a global health threat [44-47]. Nevertheless, factor in regulating the environmental

79 pathways of antibiotic resistance, especially the biotic factors, have not yet been

80 directly addressed. This hampers our ability to predict changes and risks in antibiotic

81 resistance under anthropogenic activities and global environmental change.

82 The main objective of this study was to investigate if, as proposed by the theory of

83 biological invasion, soil microbial species diversity could act as a biological barrier

84 preventing invasion by antibiotic resistance. In other words, we addressed the

85 consequences of soil diversity loss on the fate of antibiotic resistance genes in the soil

86 environment. Pig manure was used as the source of ARGs, and microbial diversity was

87 manipulated by employing a dilution–reinoculation approach (i.e. inoculating sterile

88 soil microcosms with serial dilutions of a soil microbial suspension) (Figure S1). To

89 test our hypothesis, we characterized bacterial communities and antibiotic resistome

90 using Illumina Miseq profiling of 16S ribosomal genes and high-capacity quantitative

91 PCR arrays with 296 primer sets targeting almost all major classes of ARGs [48, 49]. Our

92 study provides empirical evidence that that microbial diversity negatively relates to

93 invasibility of antibiotic resistance in soil ecosystems; and further suggests that the loss

94 in microbial diversity will likely facilitate the spread of antibiotic resistance

95 **Results and Discussion**

96 **Microbial community assembly and diversity**

97 After the OTUs were classified according to the Ribosomal Database Project (RDP)

98 database [50], the soil microbial community assemble into 17 phyla (Figure 1A) and

99 dominated by *Actinobacteria*, *Firmicutes*, *Proteobacteria*, *Bacteroidetes*,

100 *Planctomycetes* and *Chloroflexi*, which are common bacterial phyla observed in soils

101 worldwide [12, 51]. *Firmicutes* were significantly depleted along with the time, which is

102 contrasted with *Actinobacteria* and *Bacteroidetes*. Compared with the impact of time,

103 minor differences in the dominant bacterial phyla were observed when assessing

104 differences among the dilution depths. Nevertheless, *Proteobacteria* are significantly

105 ($P < 0.01$) enriched after high dilution.

106 Rarefaction curves were constructed for each individual sample showing the number of

107 observed OTUs (Figure 1B), defined at a 97% sequence similarity cut-off in QIIME [52,

108 53]. To assess the sequencing depth, we calculated Good's coverage scores (Figure 1B)

109 and they were highly comparable for all samples ranging from 94.25 to 98.51%

110 indicating that the sequencing depth was adequate to reliably describe the bacterial

111 microbiome. As expected, rarefaction curve confirmed that the dilution treatment had

112 resulted in progressively decreasing species richness and diversity, which indicated that

113 dilution-to-extinction method applied here is one of the few available methods to

114 manipulate microbial biodiversity of complex natural ecosystems such as the soil [38].

115   Furthermore, the impacts of incubation time exhibited a higher degree of variation in

116   the shape of their rarefaction curves as compared to the dilution treatments. The

117   observed OTUs (richness) and diversity of OTU were significantly depleted with the

118   increasing incubation time (Figure S2). Similarly, principal coordinates (PCoA)

119   analysis of weighted UniFrac distances between samples revealed (Figure S3) that

120   microbial communities clustering together according to their initial dilution levels

121   along with PC2 which explained 10.41% of the total variations. Whereas microbial

122   communities of different incubation time are separated along with PC1 (explained

123   53.55% of the total variations). The bacterial abundance was also measured using Real-

124   time Quantitative PCR (qPCR) based on the 16S rRNA gene. The results showed that

125   in the first ten days, the bacterial abundance was decreased in treatment of S and D0

126   (which had a relatively higher diverse native microbiota), whereas, to a lower extent, a

127   slighted increase was observed in rest of treatments (which had a relatively lower

128   diverse native microbiota). After 10 days inoculation, the bacterial abundance was

129   stable over time in all constructed microcosms, being similar between treatments. These

130   findings are consistent with those from studies showing that some of manure-derived

131   bacteria could not thrive in soil environment, and gradually decreased after manure

132   treatment, which can be attributed to the competition with resident soil bacteria and the

133   differences in environmental conditions between soil and animal gut [54-56]. Competition

134   for nutrients is an important mechanism that may limit invasions in highly diverse

135   communities [57]. Based on the resource-based niche theories, the establishment of

136   invading species is dependent on the amount of (limiting) resources that are left

137   unconsumed by native species, as well as by the rate at which native and invader species

138   consume the existing resources [58]. Additionally, most of the gut microbiota from

139   animals and humans are restricted to growth under anaerobic conditions [59], which are

140   highly different from the aerobic conditions in our microcosms. Although competition

141   for resources and shifting in oxygen condition likely affect the fate of manure-borne

142   microbiota in soil, they are only pieces of the puzzle. Other mechanisms i.e. predation

143   and negative species interactions, might also alert the survival of manure-borne species

144   [37].
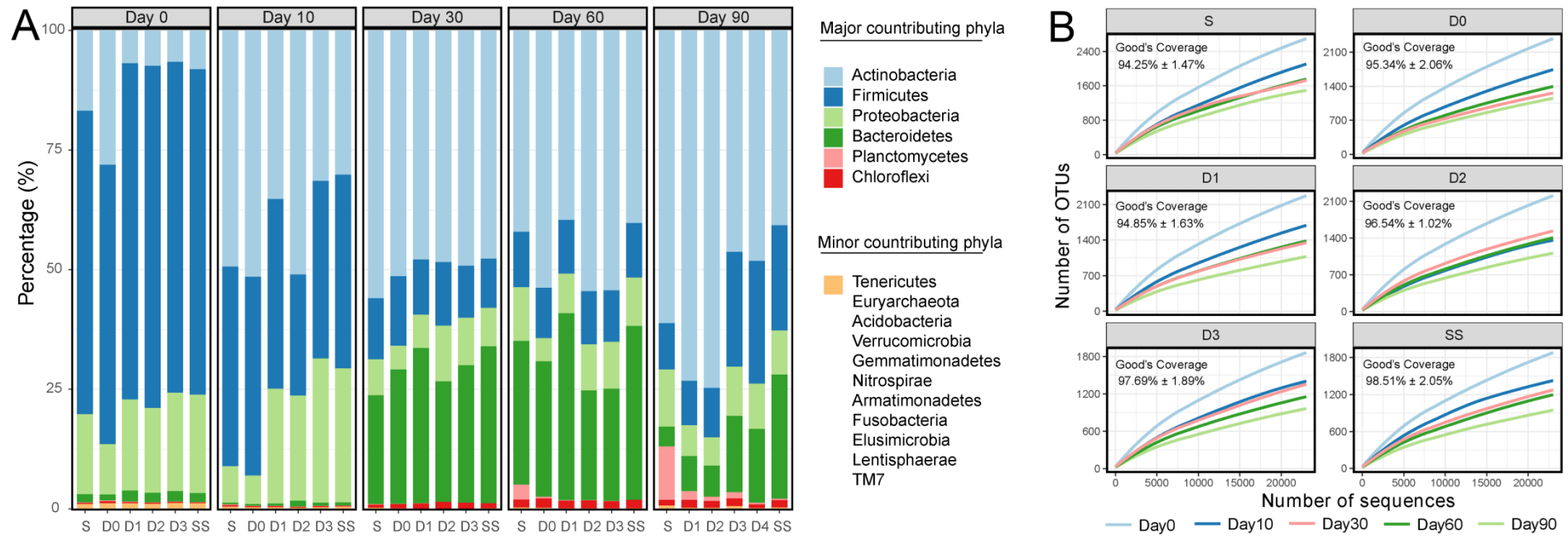
Figure 1 A: Phylum distribution of the OTUs (relative sequence abundance of bacterial phyla); B: Average Good's coverage estimates (%) and rarefaction curves of different dilution treatments. Good's coverage estimates represent averages of 20 replicates ± standard deviation.

Antibiotic resistance genes and correlation with microbial diversity

HT-qPCR was performed to investigate the abundance and diversity of ARGs. A total of 195 genes including 186 unique ARGs, class 1 integron-integrase gene and 8 transposase genes, were detected (Figure S?). The numbers of ARGs detected in each samples ranged from 47 to 116. The normalized abundance of ARG was ranging between 0.05 and 1.18 copies per 16S rRNA gene. As expected, the dilution treatment affected ARG abundance, with abundance of ARG increasing as dilution increased (Figure 2). At the begging of incubation (Day 0), because of treatments of S and D0 had a higher bacterial abundance (Figure S?), which were proposed to caused a lower normalized abundance of ARG. While the normalized abundance of ARGs between treatments became evident since day 10 and these differences were attribute to the effect of dilution, because of the bacterial abundance become stable and showed no significant different between treatments.
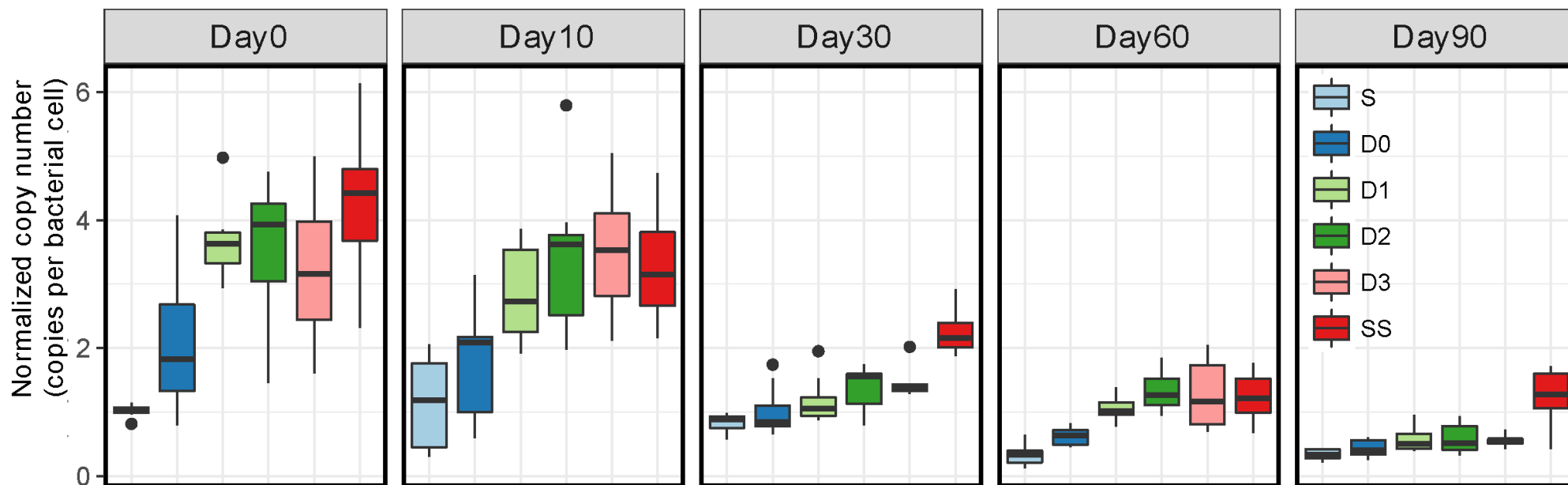
Figure 2. Normalized abundance of ARG associated with the six dilution treatments during the 90 days incubation. Box plots display the first (25%) and third (75%) quartiles, the median and the maximum and minimum observed values within each data set.

The biggest obstacle to understanding of the importance of microbial biodiversity for the functioning of ecosystems is the lack of sound experimental approaches to make directed and predictable changes in the diversity of microbial communities in soil [38]. Both the present and prior studies demonstrated that dilution-to-extinction method was an effective way to manipulate the soil microbial community to achieve a gradient in microbial diversity and species richness [30, 37]. By using ordinary least-squares (OLS) regression model, to our knowledge, we first explored the relationship between microbial diversity, estimated with the Inverse Simpson diversity indices and the spread of antibiotic resistance evaluated using the ARG abundance. Loss in microbial diversity ($P = 0.0003$, $R^2 = 0.1043$) were linearly associated with increase in ARG abundance (Figure 3). We also found biodiversity components such as phylogenetical diversity ($P = 0.0003$, $R^2 = 0.0946$) and species richness ($P = 0.0016$, $R^2 = 0.0810$) were also highly and negatively related to the dissemination of antibiotic resistance. In contrast, nonsignificant relationship between evenness and ARG abundance was found ($P = 0.8122$, $R^2 = 0.0005$). Further analyses provided evidence that Inverse Simpson diversity was positively and strongly related to phylogenetical diversity ($P < 0.0001$, $R^2 = 0.7227$) and species richness ($P < 0.0001$, $R^2 = 0.4264$). Additionally, phylogenetical diversity was also positively related to the species richness, whereas, evenness, was an exception, had no significant correlation with Inverse Simpson diversity or any of biodiversity components (Figure s?). Albeit our results are correlative in nature, and thus, the results reported here are cannot be taken as a definitive proof of causation, nevertheless, these findings are consistent with those from

theoretical and experimental studies showing that a higher diverse soil microbial communities can act as a biological barrier against invasion [32, 39, 60, 61] . For example, a recent study showed that a significant negative correlation was detected between the survival rate of *Listeria monocytogenes* L9 and the Inverse Simpson metric ($\rho = -0.817$, $P < 0.05$) and suggested that erosion of microbial diversity may have damaging effects regarding circulation of pathogenic microorganisms in the soil environment [39]. Our results provide the first empirical evidence at gene-level to show that natural ecosystems could show variable resistance to invasion by aliens, and this resistance was relate to the species diversity in the system. Finally, our results support the hypothesis that microbial diversity is the foundation for the maintenance of ecosystems [1]. Although, functional redundancy in soil microorganisms are considered prevalent, and has been thought to overwhelm any type of diversity–function relationship, i.e. initial loss in microbial diversity was unlikely to substantially affect ecosystem functions [19, 37]. However, if we take the biological barrier effect of microbial diversity into consideration, the functional redundancy could be overestimated in the previous studies.
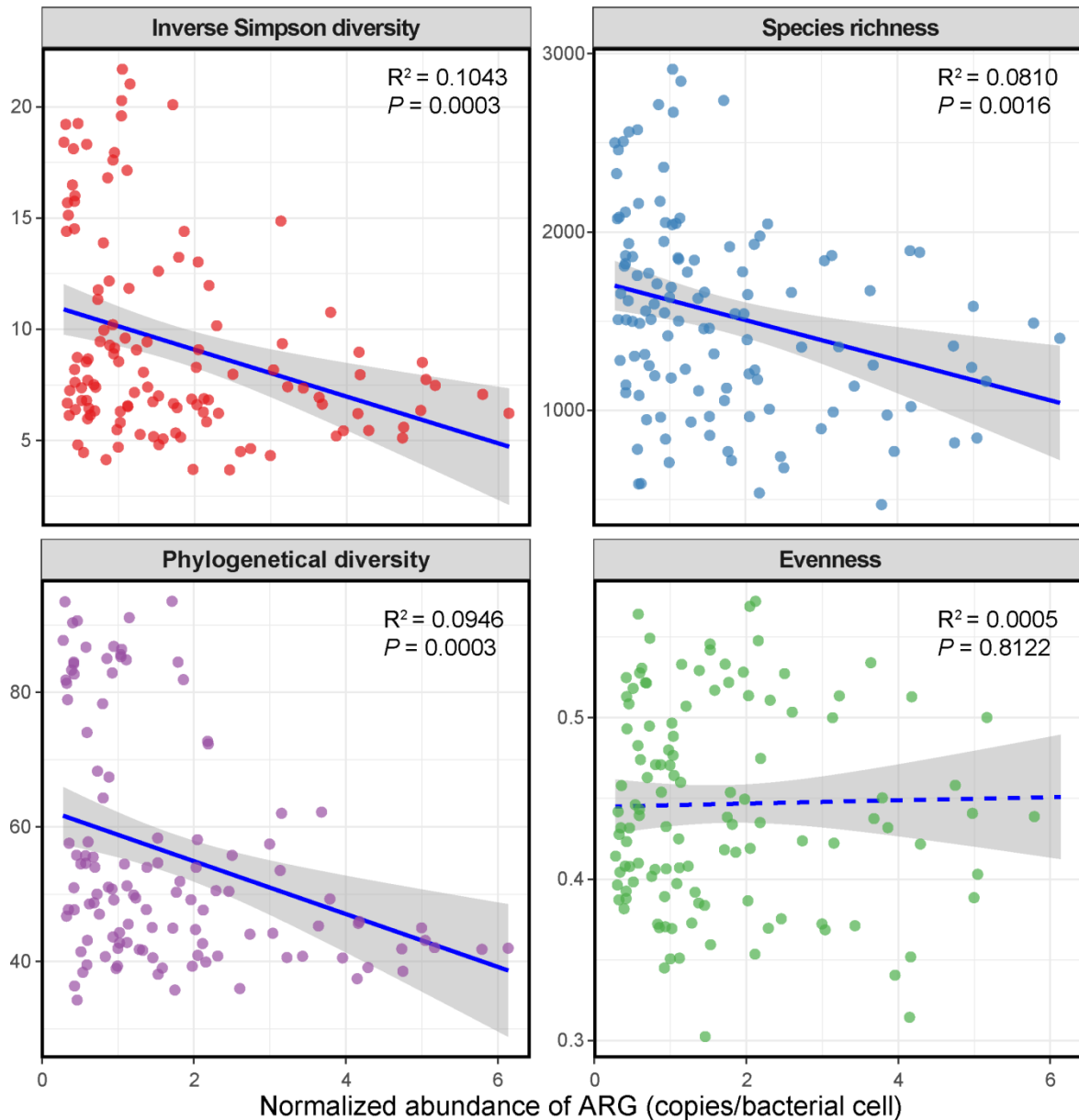
Figure 3 Ordinary least squares (OLS) regression model showing the relationships between ARG abundance, and microbial diversity, biodiversity components and evenness. The solid blue lines indicate statistical significance for the relationships, while the dashed lines indicate no statistical significance for the relationships. The shaded areas show the 95% confidence interval of the fit.
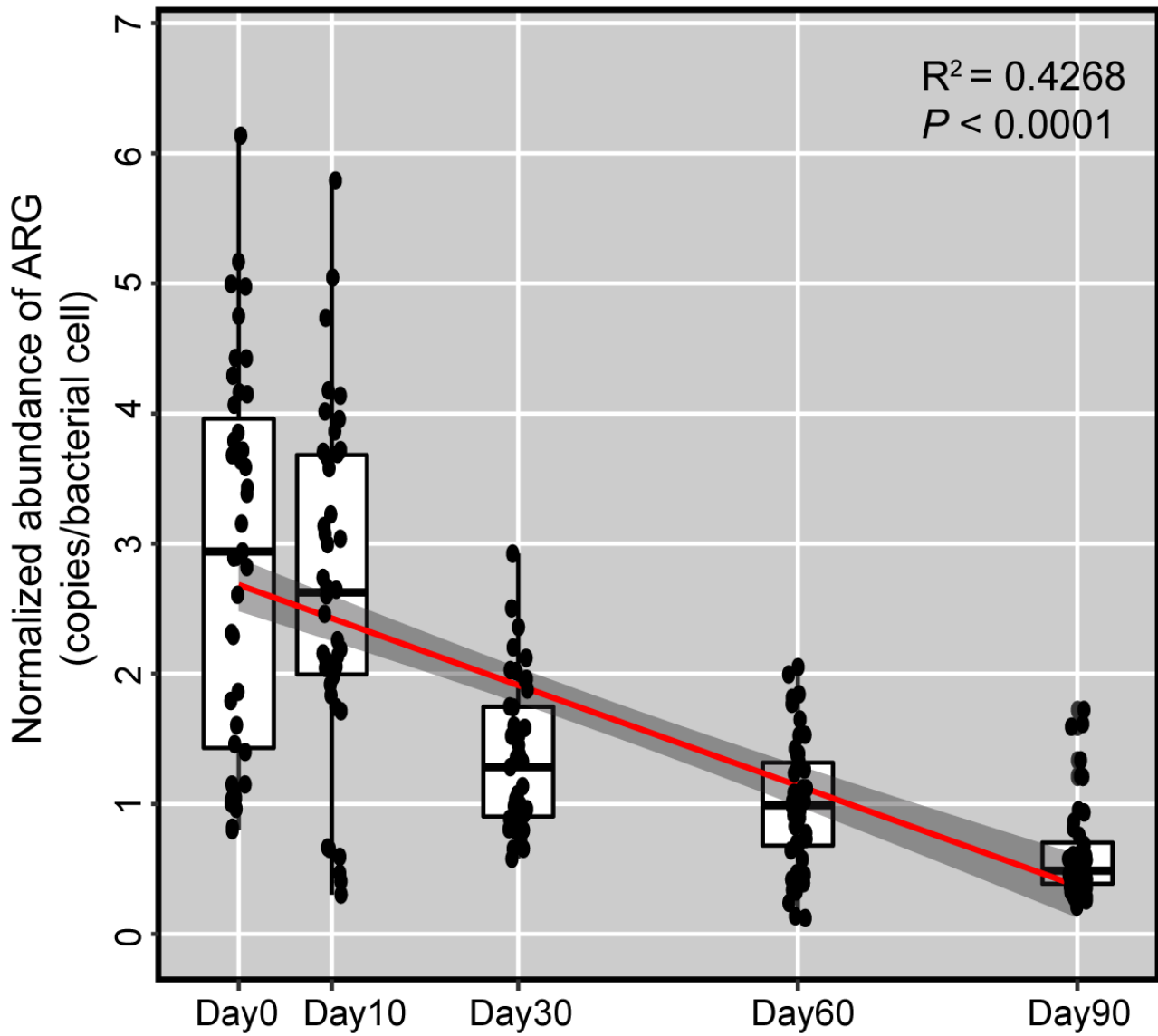
Figure 4. Relationships between ARG abundance and incubation time. The solid lines represent the fitted OLS model and the shaded areas show the 95% confidence interval of the fit. Box plots display the first (25%) and third (75%) quartiles, the median and the maximum and minimum observed values within each data set (n=24).

In addition to the impact of dilution treatment on ARG abundance, we also found that ARG abundance depleted with the increasing of incubation time (P<0.0001) (Figure 4). To further investigate the direct and indirect effects of dilution treatment and incubation time on profile of ARG, we generated structural equation models (SEMs) based on the known effects and relationships. Besides, SEM is an a priori approach offering the ability to separate multiple pathways of influence and take them as a system, and is useful to explore the complex networks of relationships found in ecosystems [62]. Except for bacterial diversity and abundance (16S rRNA copies), MGEs was also included in our models, given their effects on profiles of ARG. Our model explained 86% of the variance found in the patterns of ARGs. Incubation time and dilution treatment can directly impact the patterns of ARGs or indirectly by strongly affecting the diversity and abundance of bacteria and abundance of MGEs. Bacterial diversity posed significant directly effect on ARG (ρ=0.09, $P < 0.05$), and indirectly impacted the patterns of ARGs by strongly affecting the MGEs abundance (ρ=-0.07, $P < 0.05$). While bacterial abundance on the other hand showed non-significant directly impacts on ARG (ρ=0.02, $P > 0.05$). Despite these results, the standardized effects from SEMs revealed that MGEs abundance had a direct positive effect on the pattern of ARG and was the major contributors to this model, which was align with field studies [63], indicating that MGEs were the most dominant factor altering the ARG profiles. Both the direct and indirect effects of dilution treatment showed a positive correlation with ARG abundance, and the indirect effect (through affecting the diversity and abundance of bacteria and

abundance of MGEs) contributing larger than the direct effect. The incubation time had a negative effect on the pattern of ARG and the decrease in microbial diversity and MGEs abundance were the major driver. Compared with bacterial abundance, the diversity had a higher impacts in shaping the ARG profiles, although these impacts was much smaller than MGEs, dilution treatment and incubation time.
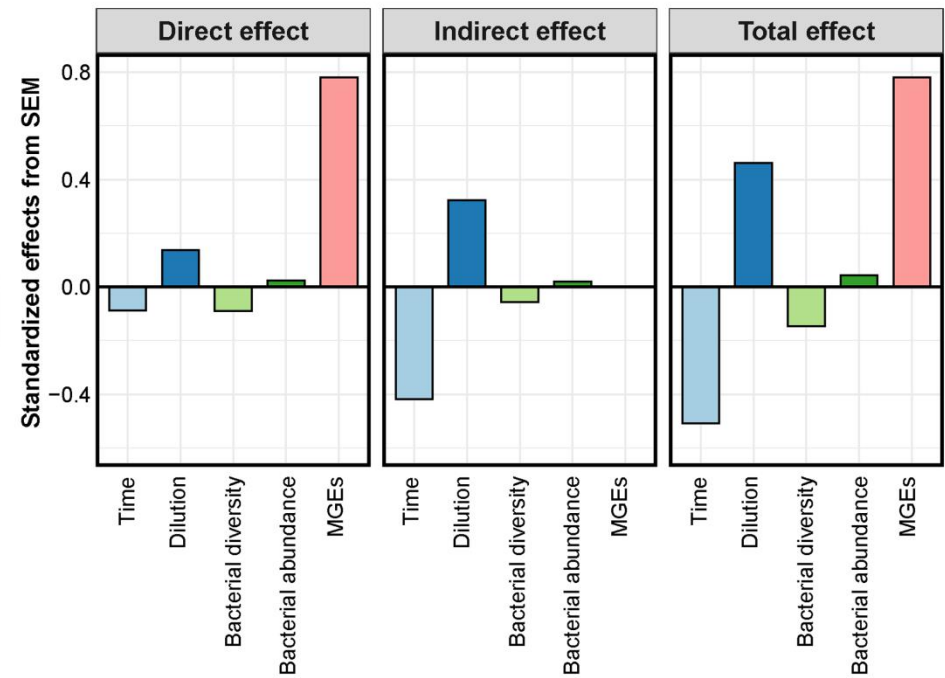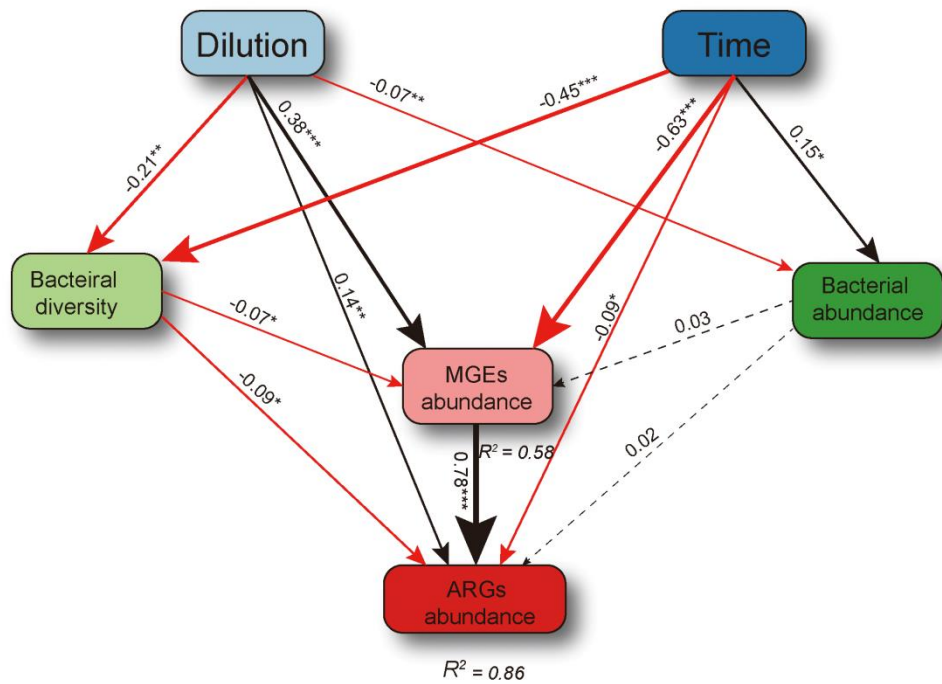
Figure 5. Structural equation models showing the direct and indirect effects of dilution treatment, time, bacterial abundance, bacterial diversity, and MGEs on the ARG patterns. Black and red arrows indicate positive and negative relationships, respectively. Continuous and dashed arrows indicate significant and nonsignificant relationships, respectively. Numbers adjacent to arrows are path coefficients, and width of the arrows is proportional to the strength of path coefficients. R^2 denotes the proportion of variance explained. Significance levels are indicated: *$P < 0.05$, **$P < 0.01$, and ***$P < 0.001$. Standardized effects (total ,direct, and ndirect effects) derived from the structural equation models. The hypothetical models fit our data well, as suggested by $\chi^2 = 0.49$, $P = 0.48$, df=1, GFI =0.99, and RMSEA = 0.00.

Conclusions

The provisioning of ecosystem services essential for human development heavily relies on the diversity of soil microorganisms. By using the microcosm model, our results provide empirical evidence that loss in microbial diversity will likely facilitate the proliferation and spread of antibiotic resistance. Our results fill a critical gap in our understanding the role of microbial diversity and provide additional insights that soil microbial communities can act as a biological barrier against invasion and the erosion of diversity may alert the circulation of antibiotic resistance genes in the environment. Altogether, ecosystems might have a limited buffering capacity of multiple ecosystem functions against biodiversity loss.

Materials and Methods

Microcosms and Experimental Setup

Surface soil (0 to 20 cm) was collected from a cropland used for planting rice in Jiaxing, Zhejiang, south China (30°50′7.7″ N, 120°43′5.7″ E). Pig manure was obtained from a local commercial pig farm. Soil and pig manure properties are described in Table S1. The soil was sieved to < 2mm, part of the soil was sterilized by γ irradiation (35 kGy), and then 3600g irradiated soil were divided into 120 flasks (200-ml) as matrix of soil microcosms, whereas the rest of soil was used, in the dilution-to-extinction experiment, as an inoculum. To test the sterility 0.5 g of irradiated soil was spreading onto tryptic soy broth (TSB) and potatodextrose agar (PDA) media []. No bacterial and fungal growth on agar plates was observed after six days. An initial soil suspension was prepared by mixing 30g soil with 50 ml autoclaved demineralized water using a ultraviolet-sterilized blender at the maximum speed for 5 minutes[], equivalent 0.6g soil/ml approximately. Totally, four levels of dilution of the soil suspension were used as inocula to create a diversity gradient of soil microorganisms ranging from undiluted (D0) to diluted 10^{-2} (D1), 10^{-4} (D2) and 10^{-8} (D3) suspensions, and 5ml of suspension was subsequently inoculated into 200 ml flasks containing 30 g dry of sterile soil, equivalent to 10^{-1}, 10^{-3}, 10^{-5}, 10^{-9} g of non-sterile soil/g sterile soil. Additionally, a positive control and negative control by using non-sterile soil (S) and sterile soil (SS) were also performed. For each treatment, four replicates were established. At last, 1.2 g pig manure as the source of antibiotic resistance was added to each flask. The flasks were then closed with sterile lids and the inoculated microcosms incubated at 20 °C,

moisture content was maintained at 70% water-holding capacity (WHC), by addition of autoclaved demineralized water. At days 10, 20, 30, 60, and 90 four replicates of each combination of soil type were collected for analysis of total bacterial abundance, diversity and composition, and determination of antibiotic resistance.

Schematic description of experimental design

Materials	Treatments		Sample collection and determinations
Unsterilized soil Sterilized soil Manure - - - - - - - - - - - - Soil suspensions Undiluted 10^{-2} 10^{-4} 10^{-8}	S	D0	Soil samples were collected after 0, 10, 20, 30, 60, and 90 days incubation. Using HT-qPCR and Illumina Sequencing to characterize the profile of antibiotic resistance gene and bacterial community.
	D1	D2	
	D3	SS	

Assessing microbial diversity

For each replicate microcosm from each dilution treatment, total DNA was extracted from 0.5g soil using a FastDNA®Spin Kit for soil according to the supplier's manual (MP Biomedical, Santa Ana, California, USA). The quality of the DNA was checked by using ND-1000 spectrophotometer (NanoDrop Technology, Wilmington, DE, USA). The concentration of DNA was determined using QubitTM dsDNA HS Assay kit through a fluorometer (Qubit^{TM} 3.0, USA). PCR was performed using 1μl of each forward (515F) and reverse (907R) bar-coded primers, 25μl of 2x ExTaq polymerase (TAKARA BIO INC, Japan) and 1μl of sample DNA as the template and 22 ul nuclease-free PCR-grade water in a total volume of 50 μl with the following PCR program of 95 °C for 5 min, followed by 30 cycles each of 95 s for 30 s, 58 °C 30 s, 72 °C for 30 s. To detect any contamination during PCR preparation, negative controls (template DNA was replaced with water) were included for all PCR reactions. PCR products of each subsample from the bar-coded primers were generated in four replicates and purified using the Wizard SV Gel and PCR Clean-Up System (TIANGEN Biotech, Beijing, China). The purified PCR products that were quantified and pooled at the same concentration, and then submitted to Illumina Hiseq2500 platform (Novogene, Beijing, China) for sequencing.

To guarantee the quality of downstream analysis, raw pair-end reads were filtered to discard raw reads containing three or more ambiguous nucleotides, or with a low ($<$ 20) average quality score, or with a short ($<$ 100 nt) length, and barcode to generate clean

joined reads capturing the complete V4-V5 region of the 16S rRNA gene by Novogene. The generated high quality sequences were processed and analyzed using QIIME pipeline []. The operational taxonomic unit (OTU) was identified using the UCLUST algorithm (Edgar, 2010) with a phylotype defined at the 97% sequence similarity level. Chimeric sequences, chloroplast and mitochondrial OTUs (around 1%), and singleton OTUs were discarded from the final OTU table. Taxonomic classification and quantification of OTUs were aligned against the Ribosomal Database Project database. Raw sequences were deposited in the National Center for Biotechnology Information (NCBI) Sequence Read Archive (SRA) under the access number SRP108158.

Assessing antibiotic resistance genes

A total of 296 primer sets were used to interrogate the soil DNA, these primers have been used, and validated in a previous study, which targeted resistance for all major classes of antibiotics (285 primer sets), transposase genes (8 primer sets), the universal class 1 integron-integrase gene (*intI1*), the clinical class 1 integron-integrase gene (*cintI1*) and the 16S rRNA gene (Supplementary Data 2). Before amplification with Wafergen SmartChip Real-time PCR system, all soil DNA was diluted to 50 ng μl^{-1} using sterile water. Amplification was conducted in a 100 nL reaction system and all qPCR reactions were conducted in triplicate and for each primer set, a non-template negative control was included. A more detailed description about the experimental procedure can be found in previous studies.

HT-qPCR data was analyzed using SmartChip qPCR software (V 2.7.0.1). Reactions with poor melting curve analysis and reactions with amplification efficiency beyond

the range (90%-110%) were discarded. Then screened with conditions that (1) a threshold cycle (CT) must be < 29 and (2) positive samples should have three replicates simultaneously. Relative copy number was calculated according to methods described in ref. 43: relative gene copy number $=10^{(29-C_T)/(10/3)}$, where C_T refers to quantitative PCR results and 29 refers to the detection limit.

To differentiate the variations in the bacterial abundance, absolute 16S rRNA copy numbers were quantified by the standard curve (SC) method with Roche 480 system (Roche Inc., USA). Each 20 µl qPCR mixture consisted of 10 µl 2× LightCycle 480 SYBR Green I Master (Roche Applied Sciences), 0.5 µg µl^{-1} bovine serum albumin (BSA), 1 µM each primer, 1 ng µl^{-1} DNA as template and 6 µl nuclease-free PCR-grade water. The thermal cycle was set according to the previous descriptions (). A plasmid control containing a cloned and sequenced 16S rRNA gene fragment $(1.18 \times 10^{10}$ copies per microlitre) was used to generate eight-point calibration curves from ten-fold dilutions for standard calculation. All qPCRs were performed in technical triplicates with non-template negative controls.

Statistical analysis

By using ordinary least squares (OLS) regression models we explored the relationships between pattern of antibiotic resistance and microbial diversity estimated with the Inverse Simpson diversity, evenness and biodiversity components including phylogenetical diversity and species richness. OLS regression was conducted with R and visualized with package of "ggplot 2". We used Structural equation model (SEM) (ref. 32) to evaluate the direct and indirect relationships between dilution treatments,

incubation time, bacterial diversity, bacterial abundance, MGEs abundance, and pattern of antibiotic resistance. SEM is an a priori approach offering the capacity to visualize the casual relationships between variables by fitting data to the models representing causal hypotheses. Thus, the first step in SEM requires establishing an a priori model based on the known effects and relationships among the drivers shifting antibiotic resistance. The theoretical model assumptions were as follows (Figure S?): (i) Dilution treatment and incubation time might have direct influences on MGEs and the ARG patterns; (ii) and also they could indirectly influence the patterns of MGEs and ARGs by changing the bacterial abundance and diversity; (iii) bacterial abundance and diversity might have direct influences on ARG patterns and they could indirectly influence the patterns of ARGs by changing the MGEs abundance and detected number. Before modeling, we examined the distributions of all of our variables and tested their normality. All data was standardized and using Z-score in the downstream analysis. Bivariate correlation was performed to examine the pairwise correlations among these variables using SPSS 20 (IBM, Armonk, NY, USA), and the covariance matrix was imported into AMOS 21 (SPSS Inc., Chicago, IL, USA) for SEMs construction using the maximum-likelihood estimation (MLE) method. We parameterized our model using our dataset and tested its overall goodness of fit. There is no single universally accepted test of overall goodness of fit for SEMs, so we used multiple goodness-of-fit criteria. We used the chi-square (χ^2) test (the model has a good fit when χ^2 is low and P - value is high, traditionally $P > 0.05$), the root mean square error of approximation (RMSEA; the model has a good fit when RMSEA is near 0 and the probability is high, traditionally

P > 0.05), high goodness-of-fit index (> 0.90) and low Akaike information criteria (AIC). With a reasonable model fit we were free to interpret the path coefficients of the model and their associated *P*-values. A path coefficient is analogous to a partial correlation coefficient and describes the strength and sign of the relationships between two variables. The solid and dot line represent a significant (*P* < 0.05) and nonsignificant correlation (*P* > 0.05), respectively. And we used black and red line to display the positive and negative correlation. Additionally, SEM is capable to partition direct and indirect effects that one variable may have on another and estimate the strengths of these multiple effects. To aid final interpretation in light of this capability of SEM, we calculated the standardized direct, indirect and total effects of dilution treatment, incubation time, bacterial abundance, bacterial diversity, and abundance of MGEs. The net effect of one factor upon another was calculated by summing all direct and indirect pathways between the two factors. If the model fits the data well, the total effect should approximate the bivariate correlation coefficient for that pair of factors

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