Interactive comment on “Ammonium and nitrate additions differentially affect soil microbial biomass of different communities and enzyme activities in slash pine plantation in subtropical China” by Chuang Zhang et al.

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Comments in response to Referee 2
This manuscript by Zhang et al examines the impact of 4 years of simulated atmospheric N deposition in 30 year old pine-dominated forest plots. The approach used by the authors is novel in that it differentiates between type of N applied (NH4 vs NO3), which is particularly interesting due to the global stabilization/decline in N deposition as NO3 and the increase in N deposition as NH4.

Response: We would like to thank you for the helpful and constructive comments, which further improved the manuscript. We have carefully revised our manuscript to take account of your comments and suggestions. Please find below our responses to comments. The page and line numbers mentioned here refer to the latest revision of our manuscript simultaneously submitted with all figures as a single PDF file.

General Comments:
1) In general, the data are much more complex across time than the authors present. It would be nice if the results are as clean as suggested in the topic sentence of each discussion paragraph, but it is simply not the case because many of the results are time-dependent. For this reason, the authors need to greatly expand the interpretation and discussion of the treatment x time interaction that is presented in Table 2. Further, to help the reader reason through the data, I think it would be beneficial to collapse the data to not include the 3 sampling times for those factors that do not exhibit a significant treatment x time interaction. For example, Fig 2(i) can be reduced to three bars for control, ammonium N, and nitrate N because there was not a significant interaction.

Response: We discussed the possible reasons resulted in the interaction effects at the discussion section. “The N treatments also varied significantly on a seasonal basis and there were interaction effects between N treatments and seasons on the contents of some PLFA biomarkers and enzyme activities (Table 2). Climate conditions, plant growth, the amount of litter returned, and plant-soil-microorganism systems varied across the three seasons. The temperature ranged from 13.5 to 27.6 °C, and precipitation ranged from 88.2 to 176.6 mm, across the three seasons (Fig. S1), and did not limit the growth of microorganisms. The positive relationships between PLFA biomarker contents and soil moisture contents indicate that soil moisture had a strong influence on soil microbial community biomass. There may be interaction effects between plant growth, the mass and quality of litter, plant-microbe competition, and soil

C1

C2
nutrient dynamics. For example, compared with the control plots, the soil DOC contents were lower, and soil nitrate contents stayed the same in June (the growing season) in the ammonium treatment, but the soil DOC and nitrate contents were higher in the ammonium and nitrate treatments in March and October (the non-growing season, Fig. 2). This indicates that there was stronger competition between plants and microbes for available C and N in June than in March and October, and that there were interaction effects between plants and microbes on soil C and N availability. This might explain the interaction effects between N additions and seasons on the activities of C and N-acquisition enzymes. The effects of interactions between N additions and season on the AMF PLFA contents, along with available C and N dynamics, may result from plant growth as plant-AMF symbiotic systems may be influenced by fine root biomass.” (line 354-371)

We also simplified the figure that the indexes were treatments-independent, and the figures were shown at Fig. 1 and Fig. 3.

2) More can be done with soil enzyme data to forward the authors main hypotheses and ideas that are introduced in line 119. For example, enzyme data can be presented as ratios of C acquiring/N acquiring and/or C acquiring/P acquiring. Such analysis will provide a clearer avenue to draw conclusions about whether microbes can alter how resources are allocated to scavenge for nutrients under different conditions.

Response: “We compared the stoichiometry of C and P to N-acquisition enzyme activities by ln(aG+BG+CBH+BX) and lnAP to lnNAG, respectively (n=27).” (line 210-212) “When compared to control, the ratios of C to N-acquisition enzyme activities were about 0.2 higher, the ratios of N to P acquisition enzyme activities were about 0.1 lower, and there were no obvious differences in the ratios of C to P acquisition enzyme activities in the ammonium and nitrate treatments.” (line 277-280) “The ratios of C or P to N acquisition enzyme activities were higher in the ammonium and nitrate treatments than in the control plots, and the N-acquisition enzyme activities per unit of microbial biomass were lower in the ammonium and nitrate treatments than in the control (Fig. 5), indicating that microorganisms secreted enzymes in line with the economic theory. Measured absolute enzyme activities were positively correlated with soil pH and ammonium contents, and negatively correlated with nitrate contents (Fig. 6). The inhibitory effects of N on the soil absolute enzyme activities may be more closely related to abiotic factors, i.e. soil pH and nitrification, than biotic factors (Kivlin et al., 2016).” (line 339-346)

3) Is there any ecological rationale for the March/June/October time points? What is the climatic variation across these times? Also, I assume soil moisture was measured, and if it was, those data should be presented and included in all analyses (including the RDA). Soil moisture has been shown to be a major driver of microbial community composition.

Response: “We collected soil samples in March, June, and October of 2015, to represent spring, summer, and fall.” (line 169-170) “The atmospheric conditions and plant-derived litters differed between the three seasons, and so indirectly affected the soil microbial biomass and enzyme activities of different communities. We collected soils from three seasons so that we could investigate the synthetic responses of soil microbial biomass and enzyme activities to ammonium and nitrate additions and to obtain improved information to support predictions of the effects of elevated N depositions on C, N, and P cycling.” (line 172-177) “Soil water contents (SWC) were measured by the oven drying method (105 °C).” (line 184-185) “SWC were positively correlated with soil PLFA biomarker contents, but were not correlated with the absolute enzyme activities (Fig. 6).” (line 302-303) “The temperature ranged from 13.5 to 27.6 °C, and precipitation ranged from 88.2 to 176.6 mm, across the three seasons (Fig. S1), and did not limit the growth of microorganisms. The positive relationships between PLFA biomarker contents and soil moisture contents indicate that soil moisture had a strong influence on soil microbial community biomass.” (line 357-360) The data applied to RDA analysis included soil moisture contents, and the figure was shown at Fig. 6. Average monthly atmospheric temperature and precipitation at the study site during 2015...
were shown at Fig. S1.

4) The manuscript needs to be edited for grammar, flow, and word choice. The writing is poor and must be improved significantly in order to be publishable.

Response: We have our revised version manuscript professionally edited by a native English speaker colleague, Dr Deborah Ballantine from the United International College, Beijing Normal University and Hong Kong Baptist University, Zhuhai, Guangdong Province.

Minor comments:

Line 169: Is there rationale for the dose of N applied? Any relation to predictions for future N deposition in the region?

Response: “Background atmospheric wet N deposition of about 33 kg N ha\(^{-1}\) yr\(^{-1}\) comprises 11 kg N ha\(^{-1}\) yr\(^{-1}\) as ammonium and 8 kg N ha\(^{-1}\) yr\(^{-1}\) as nitrate (Zhu et al., 2014). We established a control and test plots at the experimental sites. We equally added two types of N to the test plots, i.e. ammonium (Nammonium) as ammonium chloride (NH\(_4\)Cl) and nitrate (Nnitrate) as sodium nitrate (NaNO\(_3\)), at an annual rate of 40 kg N ha\(^{-1}\) yr\(^{-1}\). This rate was about double the background N wet deposition.” (line 154-159)

Line 229: How were the 3 sampling times considered for the RDA? Was the RDA ran on data from one of the three sampling times? Or from average data across the sampling times? Given the treatment by time interaction, this point needs clarification.

Response: The response of soil biomass of different microbial communities and enzyme activities to N treatments was similar in the three sampling seasons, so all of data in the treatments and seasons (n=27) was applied to RDA analysis. And we added the n value to statistical analyses section. (line 231)

Line 315: Others have shown that N addition disproportionately effects soil fungi and may stimulate soil bacteria (for example, see doi 10.3389/fmicb.2016.00259 and 10.1128/AEM.01224-14). This dynamic may also help explain the increase in DOC observed with N addition.

Response: We have added that “Moreover, the higher soil DOC concentrations observed in the nitrate-addition treatments (Fig. 2) may be attributed to changes in the diversity of the composition of saprophytic bacteria (Freedman and Zak, 2014; Freedman et al., 2016).” (line 324-326)

Table 2: I think P-values can be removed from this table. Given that significant values are bolded, P-values are redundant and make the table busy for the reader.

Response: Revised as recommended, please refer to Table 1.

Please also note the supplement to this comment: https://www.biogeosciences-discuss.net/bg-2017-179/bg-2017-179-SC2-supplement.pdf

Fig. 1. The effects of ammonium and nitrate additions on soil pH and ammonium contents. Small letters represent significant differences between treatments \((P < 0.05)\), and error bars represent means ± standard errors \((n=9)\).

Fig. 2. The effects of ammonium and nitrate additions on soil nitrate and DOC contents for each sampling event. Capital letters represent significant differences between the treatments \((P < 0.05)\), and small letters represent significant differences between the sampling events \((P < 0.05)\). Error bars represent means ± standard errors \((n=3)\).
Fig. 3. The effects of ammonium and nitrate additions on Total PLFAs, PLFA contents of bacteria, G− and G+/G−. Small letters represent significant differences between treatments ($P < 0.05$), error bars represent means ± standard errors (n=9). G+ represents gram positive bacteria and G− represents gram negative bacteria.

Fig. 4. The effects of ammonium and nitrate additions on PLFA contents of fungi, actinomycetes, AMF, SAP, G+, and fungi/bacteria ratio for each sampling event. Capital letters represent significant differences between the treatments ($P < 0.05$), and small letters represent significant differences between the sampling time ($P < 0.05$), error bars represent means ± standard errors (n=3). G+ is gram positive bacteria, AMF is arbuscular mycorrhizal fungi, and SAP is saprophytic fungi.
Fig. 5. The effects of ammonium and nitrate additions on N, P-acquisition specific enzyme activities for each sampling event. Capital letters represent significant differences between the treatments ($P<0.05$), and small letters represent significant differences between the sampling time ($P<0.05$), error bars represent means ± standard errors (n=3).

Fig. 6. Redundancy analyses between (a) soil properties and enzyme activities, and (b) soil properties and PLFA-biomarker contents.