

Nitrogen availability influences natural abundance ^{15}N of *Aspergillus oryzae*

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Nitrogen availability controls nitrogen mineralization and nitrification which are important reaction for nitrogen cycle in the soil (Schimel and Bennet. 2004). To evaluate nitrogen availability, soil C/N ratio and net nitrogen mineralization are usually used. However, difficulty in extracting the nitrogen source pools for soil microbes and getting the field circumstance information by using laboratory culture experiment create the difficulty in evaluating the nitrogen availability accurately. Then, the natural abundance of ^{15}N ($\delta^{15}\text{N}$) has been used for evaluating the nitrogen availability as a tool of getting the field circumstance information. Dijkstra et al. (2008) showed negative correlation between $\delta^{15}\text{N}$ which means the difference between $\delta^{15}\text{N}$ of SMB (Soil Microbial Biomass) and $\delta^{15}\text{N}$ of microbial substrate (K_2SO_4 extractable nitrogen from soil) and microbial substrate C/N, and this result suggested $\delta^{15}\text{N}$ could be a good indicator for nitrogen availability. They explained this phenomenon that mineralization is the dominant process for soil microbes at the high nitrogen availability sites, and SMB becomes enriched in ^{15}N because microbes release NH_4^+ which is depleted in ^{15}N . However, previous study about the relationship between $\delta^{15}\text{N}$ -biomass and $\delta^{15}\text{N}$ - NH_4^+ in C/N controlled pure culture is conducted only by Collins et al. (2008) who used *E. coli*, and they could not detect $\delta^{15}\text{N}$ - NH_4^+ in a low concentration. Thus, the relationship between nitrogen availability and $\delta^{15}\text{N}$ -biomass is unclear. The purpose of our study is to reevaluate if biomass becomes enriched in ^{15}N when microbes release NH_4^+ which is depleted in ^{15}N . In this study, we cultured Fungi (*Aspergillus oryzae*) who has large biomass in the forest soil in C/N controlled pure culture (C/N5, 10, 30, 50, 100) for 4 days. We used glycine and glucose as a nitrogen and carbon source. And we measured mainly changes in $\delta^{15}\text{N}$ -biomass, NH_4^+ concentration and $\delta^{15}\text{N}$ - NH_4^+ . In C/N5 and 10 where NH_4^+ concentration increased over time, we found that biomass was strongly enriched in ^{15}N and NH_4^+ is depleted in ^{15}N . Conversely, in C/N 30, 50 and 100 where microbes hardly released NH_4^+ , we found that $\delta^{15}\text{N}$ -biomass got the almost same value of initial $\delta^{15}\text{N}$ -glycine. In the presentation, we will discuss more detail about the carbon and nitrogen mass balance during our experiment.