

南極コケ坊主生態系における窒素固定菌と脱窒菌の多様性

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Microfloral association for nitrogen cycling in an Antarctic moss pillar inferred from phylogenetic analyses

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Aquatic mosses form unique tower-like structures called “moss pillars (Koke Bouzu in Japanese)” in ultraoligotrophic Antarctic lakes. The pillars consist of distinct redox-affected sections, that is, oxidative exteriors and reductive interiors. On the basis of analyses of fatty acid compositions, 16S/18S rRNA phylotypes, and RuBisCO genotypes (Nakai et al. *Polar Biology* 2012a, 2012b, 2012c), we had proposed that a “pillar” is a community and habitat of functionally interdependent organisms. Here, we report the diversity of the genes encoding nitrogenase (*nifH*), nitrite reductase (*nirK* and *nirS*), and nitric oxide reductase (*qnorB*) in a pillar. In total, 56 PCR clone libraries were constructed from an entire pillar, and 96 clones from each library (a total of 5,376 clones) were sequenced. Phylogenetic analyses showed that the *nifH* gene sequences of purple photosynthetic γ -proteobacteria, the *nirK* sequences of α -proteobacteria, the *nirS* sequences of β -proteobacteria, and the putative *qnorB* sequences of acidobacteria were dominant in the pillar. Furthermore, cyanobacterial *nifH* sequences were detected only in the exterior of the pillar, whereas sulfate-reducing δ -proteobacterial *nifH* sequences were subdominant in the interior. Such layer-specific distributions were also found during *nirK*, *nirS*, and *qnorB* sequence analyses. These results suggest that different phylogenetic groups participate in the nitrogen fixation and denitrification processes within a pillar. We would like to discuss about the microbial synergy that plays a vital role in the existence and maintenance of the moss pillar ecosystem.

References

1. Nakai R, Abe T, Baba T, Imura S, Kagoshima H, Kanda H, Kanekiyo A, Kohara Y, Koi A, Nakamura K, Narita T, Niki H, Yanagihara K, Naganuma T (2012a) Microflorae of aquatic moss pillars in a freshwater lake, East Antarctica, based on fatty acid and 16S rRNA gene analyses. *Polar Biology* **35**(3): 425–433.
2. Nakai R, Abe T, Baba T, Imura S, Kagoshima H, Kanda H, Kohara Y, Koi A, Niki H, Yanagihara K, Naganuma T (2012b) Eukaryotic phylotypes in aquatic moss pillars inhabiting a freshwater lake in East Antarctica, based on 18S rRNA gene analysis. *Polar Biology* **35**(10): 1495–1504.
3. Nakai R, Abe T, Baba T, Imura S, Kagoshima H, Kanda H, Kohara Y, Koi A, Niki H, Yanagihara K, Naganuma T (2012c) Diversity of RuBisCO gene responsible for CO₂ fixation in an Antarctic moss pillar. *Polar Biology* Published online before print (doi:10.1007/s00300-012-1204-5)