

3.101 Variations in atmospheric CO₂ and its $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ observed at Minamitorishima Island in the western North Pacific.

Presenting Author:

Shohei Murayama, National Institute of Advanced Industrial Science and Technology (AIST), Tsukuba, Ibaraki, Japan, s.murayama@aist.go.jp

Co-Authors:

Kazuhiro Tsuboi, Meteorological Research Institute, Tsukuba, Ibaraki, Japan

Shigeyuki Ishidoya, National Institute of Advanced Industrial Science and Technology (AIST), Tsukuba, Ibaraki, Japan

Hidekazu Matsueda, Meteorological Research Institute, Tsukuba, Ibaraki, Japan

Yosuke Niwa, Meteorological Research Institute, Tsukuba, Ibaraki, Japan

Yosuke Sawa, Meteorological Research Institute, Tsukuba, Ibaraki, Japan

Shinya Takatsuji, Japan Meteorological Agency, Chiyoda-ku, Tokyo, Japan

Kazuyuki Saito, Japan Meteorological Agency, Chiyoda-ku, Tokyo, Japan

Kentaro Kozumi, Japan Meteorological Agency, Chiyoda-ku, Tokyo, Japan

Haruka Koda, Japan Meteorological Agency, Chiyoda-ku, Tokyo, Japan

Shuichi Hosokawa, Japan Meteorological Agency, Chiyoda-ku, Tokyo, Japan

Masamichi Nakamura, Japan Meteorological Agency, Chiyoda-ku, Tokyo, Japan

Tetsuyuki Usami, National Institute of Advanced Industrial Science and Technology (AIST), Tsukuba, Ibaraki, Japan

Shinji Morimoto, Center for Atmospheric and Oceanic Studies, Graduate School of Science, Tohoku University, Sendai, Miyagi, Japan

Abstract:

The station of Minamitorishima (MNM; 24°18'N, 153°58'E) is a unique monitoring site of background air at northern mid latitudes, which is situated on a remote coral island in the western North Pacific, about 1,950 km southeast of Tokyo. The Japan Meteorological Agency has made long-term continuous observations of atmospheric concentrations of major greenhouse gases at MNM. For better understandings of mechanism governing the CO₂ variation, systematic measurements of atmospheric CO₂ and its $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ have been carried out at MNM by analyzing discrete flask air samples since 2011. The CO₂ concentration shows a clear seasonal cycle with a decrease from summer to early autumn and an increase from late autumn to early spring, which is in opposite phase with the $\delta^{13}\text{C}$ cycle. Comparative analyses between both of the seasonal components suggest that the cycles are due mainly to a seasonal-dependent CO₂ exchange with C₃ plants in land biosphere. However, the variation from June to October is found to be related to CO₂ exchange with a significantly heavier $\delta^{13}\text{C}$ signal compared to the other period of the year. On the other hand, $\delta^{18}\text{O}$ shows a seasonal cycle with a decrease from summer to late autumn and an increase thereafter until early summer, due to influences of not only carbon but also hydrological cycles. Secular increase of the CO₂ concentration and decrease of $\delta^{13}\text{C}$ due to anthropogenic CO₂ emission are also observed, accompanied by year-to-year variations in opposite phase with each other,

while $\delta^{18}\text{O}$ shows a secular increase trend until 2016 and then a decrease trend. These secular trends may reflect variations in global carbon and hydrological cycles associated with the ENSO events.

In addition to these results, preliminary results of continuous measurements of the $\delta^{13}\text{C}$ and $\delta^{18}\text{O}$ using a laser spectroscopy initiated in March 2018 will also be presented.