

[JJ] Evening Poster | M (Multidisciplinary and Interdisciplinary) | M-IS Intersection

## [M-IS14]Biogeochemistry

convener: Keisuke Koba (Center for Ecological Research, Kyoto University), Hideaki Shibata (Field Science Center for Northern Biosphere, Hokkaido University), Naohiko Ohkouchi (海洋研究開発機構, 共同), Youhei Yamashita (Faculty of Environmental Earth Science, Hokkaido University)

Wed. May 23, 2018 5:15 PM - 6:30 PM Poster Hall (International Exhibition Hall7, Makuhari Messe)

Biogeochemistry is an interdisciplinary study field including ecology, geochemistry, oceanography, limnology, hydrology, soil science and environmental sciences. Respective researches have tended to be conducted separately so far. This session aims to provide a common platform for biogeochemists of different disciplines, which facilitates the interactive discussion and information exchanges for further development of biogeochemical studies.

## [MIS14-P03]Seasonal variation of O<sub>3</sub> flux in red pine forest

Junpei Ito<sup>1</sup>, \*Ryuichi Wada<sup>1</sup>, Satoru Takanashi<sup>2</sup>, Takafumi Miyama<sup>2</sup>, Michiaki Okano<sup>2</sup>, Takashi Nakano<sup>3</sup>, Tomoki Mochizuki<sup>4</sup>, Akira Tani<sup>4</sup>, Seiichiro Yonemura<sup>5</sup>, Yutaka Matsumi<sup>6</sup>, Kentaro Takagi<sup>7</sup>, Masahito Ueyama<sup>8</sup>, Yuzo Miyazaki<sup>7</sup> (1. Teikyo University of Science, 2. Forestry and Forest Products Research Institute, 3. Mount Fuji Research Institute, 4. University of Shizuoka, 5. National Agriculture and Food Research Organization, 6. Nagoya University, 7. Hokkaido University, 8. Osaka Prefecture University)

Keywords: Ozone, Forest, Flux

The emission and absorption of trace gases at the biosphere affects to atmospheric chemistry, and thus it makes influence with potential indirect effects on carbon cycle and climate (Ollinger et al., 2002). We observed O<sub>3</sub> flux with the gradient method at a meteorological tower in red pine forest (Site Code: FJY) since 2016. We also measured CO<sub>2</sub> flux at the same meteorological tower for validation of the system by comparison with CO<sub>2</sub> flux determined by the eddy covariance method. The heights of the forest canopy and the meteorological tower were about 24 m and 32 m. Concentrations of O<sub>3</sub> and CO<sub>2</sub> were measured at two heights (26 m and 34 m) above the canopy by an ultraviolet absorption O<sub>3</sub> analyzer (Thermo: 49C), and an infrared absorption CO<sub>2</sub> analyzer (Licor: LI-820). The O<sub>3</sub> instrument was calibrated once a year and the CO<sub>2</sub> instruments were calibrated every three weeks at the observation site. The air was sampled every 300 seconds from each two vertical heights and supplied to the analytical instruments through PFA tube. Concentration of CO<sub>2</sub> was also measured by an infrared absorption CO<sub>2</sub> analyzer (Licor: LI-6262) at 26.5 m to determine CO<sub>2</sub> fluxes by the eddy covariance method. Wind speed and wind direction were measured at 26.5 m and they were used to obtain fluxes by the gradient and eddy covariance methods. The CO<sub>2</sub> fluxes in the day time (9:00-16:00) in 2017 were observed with the gradient and the eddy covariance method as  $-0.14 \pm 0.12 \text{ mol m}^{-2} \text{ d}^{-1}$  and  $-0.21 \pm 0.16 \text{ mol m}^{-2} \text{ d}^{-1}$ , respectively. The CO<sub>2</sub> flux obtained by the gradient method was slightly lower and more scattered than CO<sub>2</sub> flux obtained by the eddy covariance method; however these values reasonably agreed. We made sure the flux observation system with gradient method worked properly. The observed O<sub>3</sub> concentration showed a seasonal variation and was in a maximum in May. However the primary result showed that O<sub>3</sub> deposition in the red pine forest in daytime had a maximum between June and July. The peak of O<sub>3</sub> deposition delayed to the peak of O<sub>3</sub> concentration. A chemiluminescence O<sub>3</sub> analyser was developed and optimized the measurement condition using O<sub>3</sub> standard gas generated by O<sub>3</sub> generator. The developed analyser was tested in Uenohara for 4 days and the observed O<sub>3</sub> concentrations agreed with the concentrations observed by an ultraviolet absorption O<sub>3</sub> analyzer.