

# In-situ continuous observations of atmospheric O<sub>2</sub>/N<sub>2</sub> ratio and CO<sub>2</sub> concentration on-board “SHIRASE” in the Southern Ocean

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Atmospheric O<sub>2</sub>/N<sub>2</sub> ratio (O<sub>2</sub> concentration) and CO<sub>2</sub> concentration are closely related to each other through terrestrial biospheric activities and fossil fuel combustion. However, they are independently fluctuated when their exchanges occur between the atmosphere and the ocean. For a better understanding of the global carbon cycle, atmospheric O<sub>2</sub>/N<sub>2</sub> ratio has been observed around the world since 1990s (e.g. Manning and Keeling, 2006). Until recently, most of O<sub>2</sub>/N<sub>2</sub> ratio observations have been conducted at fixed stations on land, and a few of them have been on-board vessels (e.g. Tohjima et al., 2005). In particular, O<sub>2</sub>/N<sub>2</sub> observation data are very sparse in the Southern Ocean. In this study, we have developed a continuous measurement system for atmospheric O<sub>2</sub>/N<sub>2</sub> ratio and CO<sub>2</sub> concentration and conducted in-situ observations on-board the icebreaker “SHIRASE” in the Southern Ocean on her JARE-59 cruise. The measurement system consists of a differential fuel cell oxygen analyzer (Sable Systems, Oxzilla) and a non-dispersive infrared CO<sub>2</sub> analyzer (LI-COR, LI-6252), together with a precise pressure and flow-rate control system. Repeatability of the O<sub>2</sub>/N<sub>2</sub> ratio and CO<sub>2</sub> concentration measurements was estimated to be  $\pm 4.1$  per meg (corresponding to about 0.9 ppm) and  $\pm 0.04$  ppm for 10-minute averages and  $\pm 1.6$  per meg (0.3 ppm) and  $\pm 0.02$  ppm for 1-hour averages. The measurements were started at Fremantle, Australia (FRM: 32°03'S, 115°45'E) on December 1, 2017, and stopped at Sydney, Australia (SYD: 33°52'S, 151°12'E) on March 18, 2018. Between FRM and Syowa Station (SYO: 69°00'S, 39°35'E), latitudinal gradients were observed in the O<sub>2</sub>/N<sub>2</sub> ratio and CO<sub>2</sub> concentration and their gradients showed negative correlation; the O<sub>2</sub>/N<sub>2</sub> ratio and the CO<sub>2</sub> concentration became lower and higher, respectively, towards southern latitudes. In addition, the atmospheric potential oxygen (APO  $\sim$  O<sub>2</sub> + 1.1CO<sub>2</sub>) also showed similar latitudinal gradient as the O<sub>2</sub>/N<sub>2</sub> ratio. Considering that the APO signal is mainly produced by oceanic O<sub>2</sub> flux, it is suggested that the spatial distribution of the O<sub>2</sub>/N<sub>2</sub> ratio is affected by O<sub>2</sub> outgassing from ocean surface in the southern middle latitudes. Similarly, negative correlation between the O<sub>2</sub>/N<sub>2</sub> ratio and the CO<sub>2</sub> concentration was also found in their short-term variations observed on December 9. The variation ratio of the O<sub>2</sub> concentration with respect to the CO<sub>2</sub> concentration ( $-\Delta O_2/\Delta CO_2$ ) for the short-term variation is calculated to be 5.5 ppm/ppm. This number is much larger than that expected for fossil fuel combustion (1.4 ppm/ppm) and for terrestrial biospheric activity (1.1 ppm/ppm). Since O<sub>2</sub> and CO<sub>2</sub> exchange between the atmosphere and the surface ocean occurs independently and the exchange rate is much faster in the O<sub>2</sub> exchange, it is suggested that the observed short-term variation of the O<sub>2</sub>/N<sub>2</sub> ratio is mainly produced by O<sub>2</sub> exchanges between the atmosphere and the surface ocean. In the presentation, we will show the details of our measurement system and the latitudinal and longitudinal distributions of the observed O<sub>2</sub>/N<sub>2</sub> ratio and the CO<sub>2</sub> concentration.

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## References

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