

北極域における大気ポテンシャル酸素と大気海洋間酸素フラックスの船舶観測

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Shipboard Observations of Atmospheric Potential Oxygen and air-sea O₂ flux in the northern North Pacific and the Arctic Ocean

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Simultaneous observations of Atmospheric Potential Oxygen (APO = O₂+1.1xCO₂) and air-sea O₂ flux were carried out onboard a research vessel MIRAI in the northern North Pacific and the Arctic Ocean in the autumns of 2012-2015. Air samples to measure the atmospheric δ(O₂/N₂) and CO₂ concentration to derive APO were collected on average once per day for the period September 5 – October 15, 2012, August 29 – October 6, 2013, September 1 – October 9, 2014 and August 27 – October 5, 2015. Dissolved oxygen concentration in the near-surface water was also measured continuously during these cruises, and converted to air-sea O₂ flux (F_{O₂obs}). The relationships of δ(O₂/N₂) and simultaneously-measured δ¹³C with CO₂ concentration indicated that terrestrial biospheric activities and the air-sea O₂ flux are the main contributors to the observed variations in CO₂ concentration and APO, respectively. To compare the observed APO values with those simulated using the monthly air-sea O₂ flux climatology taken from the TransCom experimental protocol (F_{O₂cli}), a simulation of APO using a 3-dimensional atmospheric transport model forced by F_{O₂cli} was also carried out. The observed APO showed larger short-term variations than the simulated APO, and F_{O₂obs} also showed larger variation than F_{O₂cli}. A simple calculation indicated that the short-term variations in APO produced by using F_{O₂obs} were comparable in magnitude to the observed, and the characteristics of the temporal variations in the observed APO were relatively well reproduced by the calculated APO. These results strongly suggest that the short-term variations seen in the observed APO is attributable to the short-term variations in the air-sea O₂ flux around the observation area. The F_{O₂obs} values were systematically higher than the F_{O₂cli} values in all cruises, with an average difference of about 0.3 μmolm⁻²s⁻¹. By uniformly mixing the sea-to-air O₂ flux of 0.3 μmolm⁻²s⁻¹ from the northern hemisphere ocean into the overlying atmosphere during the fall season, it was possible to explain the discrepancy between the observed and simulated seasonal APO cycles seen at Ny-Ålesund, Svalbard and Sendai, Japan in the fall season. These findings show that simultaneous ship observations of APO and F_{O₂obs} are useful to validate regional air-sea O₂ fluxes in detail (Ishidoya et al., submitted).

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References

Ishidoya, S. et al., Ship observations of atmospheric potential oxygen and regional air-sea O₂ flux in the Northern North Pacific and the Arctic Ocean, submitted to *Tellus B*.