

Air-ice CO₂ fluxes in the Arctic coastal area (Amundsen Gulf).

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Introduction

Sea ice covers about 7% of the Earth surface at its maximum seasonal extent. For decades sea ice was assumed to be an impermeable and inert barrier for air-sea exchange of CO₂ so that global climate models do not include CO₂ exchange between the oceans and the atmosphere in the polar region. However, uptake of atmospheric CO₂ over sea ice cover has been recently reported raising the need to further investigate pCO₂ dynamics in the marine cryosphere realm and related air-ice CO₂ fluxes.

1. Methodology and sampling site

We carried out direct measurements of pCO₂ within brines and related air-ice CO₂ fluxes (chamber method, *fig.1*) of Antarctic first year pack ice (during the “Sea Ice Mass Balance in Antarctica – SIMBA drifting station experiment, Bellingshausen Sea) and of Arctic first year fast ice (during the “Circumpolar Flaw Lead” project – CFL, Amundsen Gulf region of the Beaufort Sea).

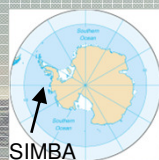
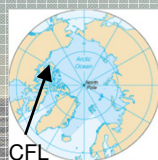


fig.1. – accumulation chamber for air-ice CO₂ fluxes measurements

2. Results (1)

Temperature exerts a strong control on pCO₂ dynamics (*fig.2*). As the temperature of the ice increases, the CO₂ in brines contained within the ice changes from a large over-saturation to a marked under – saturation (this shift occurs around -4°C). This trend is the same for both hemispheres. However, pCO₂ of the brines appears to be higher in Arctic than in Antarctica.

pCO₂ decreases because of the dilution by decaying sea ice and related decrease of DIC. In addition, dilution allows the dissolution of the carbonate minerals that promote the decrease of pCO₂.

2. Results (2)

CO₂ fluxes at air – ice interface are not detectable or weak when the ice temperature is below -6°C (*fig.3*). As the ice warms, the permeability of the ice increases, allowing CO₂ transfer at the air – ice interface.

Air-ice CO₂ fluxes are in the same order of magnitude in both hemispheres. However, in Antarctica a flooding event induced a larger heterogeneity in the ice cover and a larger variability in the CO₂ fluxes. In Arctic, the formation of superimposed ice prevented air-ice gas transfer at temperature close to 0°C.

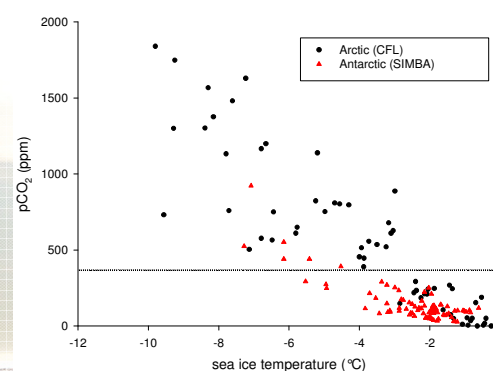


fig.2. – pCO₂ within brines versus sea ice temperature integrated over the depth of sackholes.

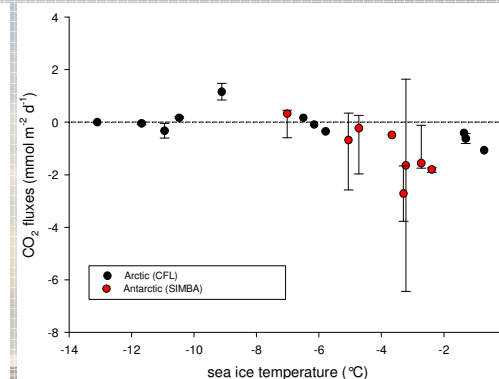


fig.3. – air-sea ice CO₂ fluxes versus sea ice temperature.

Conclusions

pCO₂ of brines exhibited high temporal variability strongly related to sea ice temperature. CO₂ fluxes also depends of the temperature of the ice for 2 reasons. First, CO₂ fluxes depend of the pCO₂ of brines that is strongly related to temperature. Second, sea ice temperature controls brine channels connections and then exchanges between brines and atmosphere. In addition, physical processes, flooding, formation of superimposed ice exert a significant influence on CO₂ fluxes.