

DECADAL CHANGES OF THE CO₂-SYSTEM PROPERTIES IN THE SUBTROPICAL SOUTH ATLANTIC: RESULTS FROM RE-OCCUPATION OF WHP A10 SECTION

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ABSTRACT

Using high-quality data for the CO₂-system and related properties obtained 10-year apart, we estimated decadal increases of anthropogenic CO₂ along the A10 section of the World Ocean Circulation Experiment (WOCE) Hydrographic Program (WHP). Increases of anthropogenic CO₂ were found down to an isopycnal surface of 27.3σ_θ (approx. 1000 dbar). In the sub-Antarctic Mode Water (SAMW), the increase was 6.9 ± 2.0 μmol kg⁻¹ on average, while in the Antarctic Intermediate Water (AAIW), it was 4.2 ± 1.9 μmol kg⁻¹. The increase in SAMW was larger in the west than that in the east of the section. No significant increases were detected in North Atlantic Deep Water (NADW) and Antarctic Bottom Water (AABW).

INTRODUCTION

In 2003, as a part of Blue Earth Global Expedition 2003 (BEAGLE), we re-visited the WHP A10 section (30°S, 50°W – 20°E) in the South Atlantic, which was occupied previously in 1992/1993. In BEAGLE, we measured properties of the CO₂-system such as surface seawater pCO₂, dissolved inorganic carbon (C_T), total alkalinity (A_T) and pH, together with dissolved oxygen, nutrients, and CFCs.

The purpose of the present study was to estimate decadal increases of anthropogenic CO₂ in the South Atlantic, and to clarify regional differences of anthropogenic CO₂ accumulation by comparing data obtained in BEAGLE with those in WHP.

OBSERVATIONS

Shipboard observations were made from November 6 to December 9 using the R/V Mirai of JAMSTEC. Atmospheric and surface seawater pCO₂ were measured continuously by a non-dispersive infrared gas analyzer. Water column C_T and A_T were measured by coulometry and potentiometry, respectively. pH was measured by spectrophotometry.

Repeatability of C_T and A_T was estimated to be 1.0 and 2.2 μmol kg⁻¹, respectively. Both C_T and A_T were set to the values of certified reference materials (batch 60) provided by Prof. A.G. Dickson of Scripps Institution of Oceanography. Repeatability of pH was 0.0010 pH unit.

APPROACH

C_T on isopycnal surfaces

To detect increases of anthropogenic CO₂ in the ocean's interior, we corrected C_T using apparent oxygen utilization (AOU) and salinity. For the correction, remineralization of organic matters was considered, while dissolution of calcium carbonate was neglected. The corrected C_T (nC_T^{CAL}) was averaged at an interval of 20° longitude on selected isopycnal surfaces, and the averaged nC_T^{CAL} in WHP was subtracted from that in BEAGLE. The *t*-Test was applied to the calculated differences (ΔnC_T^{CAL}) to check significance of decadal increases of anthropogenic CO₂ statistically.

RESULTS

Increases of anthropogenic CO₂ on isopycnal surfaces

From distributions of the ΔnC_T^{CAL} , it was found that significant increases of anthropogenic CO₂ along the A10 section were detectable down to 27.3 σ_θ (~1000 dbar). In SAMW (26.6 – 27.0 σ_θ , 350 – 700 dbar), the average of the significant ΔnC_T^{CAL} was $7.0 \pm 1.6 \mu\text{mol kg}^{-1}$, while in AAIW (27.1 – 27.5 σ_θ , 700 – 1500 dbar), it was $3.5 \pm 0.8 \mu\text{mol kg}^{-1}$. In SAMW, the increase was larger in the west than that in the east.

Increases of anthropogenic CO₂ in abyssal waters

We attempted to detect increases of anthropogenic CO₂ in abyssal waters, i.e., NADW and AABW. In NADW (salinity maximum water), no significant increases could be detected, implying that NADW does not have a measurable anthropogenic CO₂ signal yet at the latitude. In AABW, two criteria were applied to select the water: potential temperature < 1.0 °C and $\sigma_4 > 45.87 \text{ kg m}^{-3}$. From both the cases, no significant ΔnC_T^{CAL} was detected, except for 46 – 38°W by the potential temperature < 1.0 °C criterion.

CONCLUSION

From the calculated increases of anthropogenic CO₂, uptake rate of anthropogenic CO₂ was estimated to be $0.7 \pm 0.4 \text{ mol m}^{-2} \text{ yr}^{-1}$, which is close to previous estimations. This fact implies that oceanic capacity of anthropogenic CO₂ absorption did not change over the past decade.