



## Global distributions and transport of CO<sub>2</sub> in upper troposphere obtained by commercial aircraft observations

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The upper atmosphere plays an important role for global carbon cycle because it contains a large portion of atmospheric carbon as carbon dioxide (CO<sub>2</sub>) and it serves as a pathway for long-range transport. However, our knowledge about vertical structure of CO<sub>2</sub> is still limited because CO<sub>2</sub> observations are conducted generally near the surface. Measurement by using commercial airliner is very powerful tool to provide precise and global CO<sub>2</sub> distributions. Now Japan Airlines aircraft in regular service measure CO<sub>2</sub> in flight by using the Continuous CO<sub>2</sub> Measuring Equipment (CME), covering a substantial geographical area in the Comprehensive Observation Network for TRace gases by AIRLiner (CONTRAIL) project.

This study focuses on the seasonal CO<sub>2</sub> distributions observed in the upper troposphere. We used more than 2 million in-situ CO<sub>2</sub> data from 4248 flights from November 2005 to December 2009. In order to give a climatological overview of the CO<sub>2</sub> distributions in the upper atmosphere, we presented CO<sub>2</sub> for a reference year of 2008 assuming the annual trend of 1.9 ppm/year observed in recent years.

In the northern high latitudes, monthly CO<sub>2</sub> distribution in April in the upper troposphere shows larger values more than 387 ppm and values are gradually decreased to the 382 ppm in southern mid-latitude. The large decreases in CO<sub>2</sub> in the upper troposphere are detected in the northern high latitudes in boreal summer. The mixing ratios over the Eurasian continent begin to decrease in May, and it reaches minimum values of about 380 ppm in July. On the other hand, the mixing ratios over the Northern Pacific begin to decrease after one month delay and it has minimum values of 382 ppm in August. The longitudinal differences in CO<sub>2</sub> mixing ratios are largest in mid summer (~4 ppm) in July, although they are reduced to 2 ppm in August. The differences are not significant in winter season from October to April. These suggest that very low mixing ratios by strong sink for CO<sub>2</sub> in the terrestrial biosphere are efficiently transported to the upper troposphere over the Eurasian continent creating significant longitudinal differences in summer.

Next, we studied the meridional cross sections for CO<sub>2</sub> observed over the western Pacific regions in equivalent latitude-pressure coordinate. Cross sections in boreal winter to spring show clear boundaries near the equator; higher CO<sub>2</sub> mixing ratios in the Northern Hemisphere compared to lower values in the Southern Hemisphere. However, higher CO<sub>2</sub> more than 384 ppm in the Northern Hemisphere begin to get over this tropical barrier in April at higher altitudes (200-400hPa). Above 300 hPa in the Southern Hemisphere a rapid increase of about 2 ppm are found from March to June, suggesting the inter-hemispheric transport of high CO<sub>2</sub> from the Northern Hemisphere. This distribution pattern corresponds to the unique autumn maximum in the upper troposphere in the Southern Hemisphere.

CONTRAIL CO<sub>2</sub> data show the unique and clear seasonal cycles in the upper atmosphere. The longitudinal or latitudinal differences in CO<sub>2</sub> distributions are found in this frequent and wide coverage aircraft observation project. Distributions of CO<sub>2</sub> based on frequent observations delineate how CO<sub>2</sub> in the upper troposphere spreads from land to ocean or Northern Hemisphere to Southern Hemisphere.