

3.068 Measurements of carbon and hydrogen isotope ratios of atmospheric methane in the northern North Pacific and the Arctic Ocean and interpretation of Arctic methane sources.

Early Career Scientist

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Abstract:

There are large and important natural CH₄ sources in northern high latitudes, but their emissions and spatial distribution are not well understood yet. Simultaneous measurements of carbon and hydrogen isotope ratios ($\delta^{13}\text{C}$ and δD) of atmospheric CH₄ would help us to separate contributions from different types of sources (e.g., biogenic or thermogenic) to atmospheric CH₄; however, the isotope data are still sparse, especially in boreal North America and Siberia. In this study, we measured atmospheric CH₄, $\delta^{13}\text{C}$, and δD on board the research vessel MIRAI in the northern North Pacific and the Arctic Ocean in summer to autumn in 2012–2016. We also estimated the representative CH₄ isotope source signatures in their surrounding areas. A clear latitudinal gradient is observed for atmospheric CH₄, $\delta^{13}\text{C}$, and δD from 36°N to 76°N; northward increase of CH₄ and decrease of $\delta^{13}\text{C}$ and δD are evident. This suggests that biogenic CH₄ sources are dominant in northern high latitudes in the summertime. By applying a single mixing equation to the data observed at latitudes higher than 55°N, the average isotope signatures over 2012–2016 are estimated to be $-65.6 \pm 1.6\text{‰}$ for $\delta^{13}\text{C}$ and $-361 \pm 45\text{‰}$ for δD . The results are similar to the values reported previously for boreal wetland CH₄ sources. A five-day backward trajectory analysis shows that air parcels with high CH₄ come mainly from land areas of Alaska and Northern Canada and partially from Siberia. CH₄ emissions from surface water in the Arctic Ocean would not be prominent in summer to autumn in 2012–2016.