



Using atmospheric ^{14}C to provide additional constraints for global OH: results from a new approach and potential for future measurements

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The primary source of ^{14}C -containing carbon monoxide (^{14}CO) in the atmosphere is via ^{14}C production from ^{14}N by secondary cosmic rays, and the primary sink is removal by hydroxyl radicals (OH). Variations in the global abundance of ^{14}CO that are not explained by variations in ^{14}C production are mainly driven by variations in the global abundance of OH. Monitoring OH variability via methyl chloroform is becoming increasingly difficult as methyl chloroform abundance is continuing to decline. Measurements of atmospheric ^{14}CO have previously been successfully used to infer OH variability. However, these measurements have only continued at one location (Baring Head, New Zealand), which is insufficient to infer global trends.

We propose to restart global ^{14}CO monitoring with the aim of providing an additional constraint on OH variability. A new analytical system for ^{14}CO sampling and measurements has been developed, allowing for a ten-fold reduction in the required sample air volumes and simplified field logistics. The first ^{14}CO measurements from Mauna Loa Observatory show good agreement with prior measurements in the same latitude band. Preliminary work with a state-of-the-art chemical transport model is exploring sensitivity of ^{14}CO at potential sampling locations to changes in production rates and OH.

This presentation will also provide an update on a project which aims to improve the understanding of long-term OH variability via reconstructing a 150-year history of atmospheric ^{14}CO from ice cores at Law Dome, Antarctica. Sampling of the ice and on-site extractions of large volumes of ancient air were in progress during December 2018 – January 2019.