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Attribution of recent trends in atmospheric methane using inverse modelling

Joe McNorton (1), Chris Wilson (1), Manuel Gloor (2), and Martyn Chipperfield (1)

(1) Institute for Climate and Atmospheric Science, University Of Leeds, Leeds, United Kingdom (J.R.McNorton@leeds.ac.uk), (2) School of Geography, University Of Leeds, Leeds, United Kingdom

Atmospheric methane (CH₄) accounts for approximately 20% of the total direct anthropogenic radiative forcing by long-lived greenhouse gases ($0.48\pm0.05 \text{ Wm}^{-2}$), the second largest contributor after CO₂. Atmospheric observations highlight two notable changes in CH₄ since 2007. Firstly, the growth rate of methane increased to ~7ppb/yr. Secondly, the CH₄ $^{13}\text{C}/^{12}\text{C}$ -ratio ($\delta^{13}\text{C}$) has become increasingly ^{13}C -depleted. One possible explanation for both of these, is an increase in ^{13}C -depleted CH₄ emissions. This could be through increases in natural biogenic sources (e.g. wetlands), anthropogenic biogenic sources (e.g. agriculture) or a combination of both. A decrease in ^{13}C -enriched non-biogenic emissions (e.g. biomass burning) could be an explanation for the ^{13}C -depletion, but does not explain the CH₄ increase. A reduction in the atmospheric concentration of OH, the main oxidant for atmospheric methane, could also explain both ^{13}C -depletion and CH₄ increase.

We have performed a synthesis inversion using a 3-D atmospheric global chemical transport model, TOMCAT, for both CH₄ and δ^{13} C from 2005-2014. The inversion uses surface observations of both CH₄ and δ^{13} C to spatially constrain source types and possible changes to OH concentration. We will use results from this synthesis inversion to attribute the upturn in CH₄ growth to specific source and sinks, and to discuss the uncertainties in this attribution.