

A Long-Term Perspective on Recent Increases in Atmospheric CH₄ Abundance

E. Dlugokencky¹, P.M. Lang¹, K.A. Masarie¹, A. Crotwell², L. Bruhwiler¹, L. Emmons³ and P. Bergamaschi⁴

¹NOAA Earth System Research Laboratory, 325 Broadway, Boulder, CO 80305; 303-497-6228, E-mail: ed.dlugokencky@noaa.gov

²Cooperative Institute for Research in Environmental Sciences, University of Colorado, Boulder, CO 80309

³National Center for Atmospheric Research, Boulder, CO 80307

⁴European Commission Joint Research Centre, Institute for Environment and Sustainability, Ispra I-21027, Italy

After CO₂, methane (CH₄) is the most important greenhouse gas influenced by human activities. Its chemistry results in additional indirect climate effects from production of tropospheric O₃, which also affects air quality, and stratospheric H₂O. Natural emissions of CH₄ from Arctic permafrost and hydrates are susceptible to changing climate. These sources recently received attention in the media because of their potential to cause strong positive climate feedbacks. But how much of this attention is warranted?

From 1999 to 2006, the global burden of atmospheric CH₄ remained nearly constant (red line, top panel of Figure 1). During 2007 and 2008, globally averaged CH₄ increased by ~13 ppb. Do these increases signal the start of increased emissions in the Arctic from permafrost or hydrates because of warming climate? We found that the likely drivers of these increases are greater than average wetland CH₄ emissions at high northern latitudes during 2007 because of exceptionally warm temperatures and in the tropics during 2007 and 2008 because of greater than normal precipitation in wetland regions during a La Niña. These increases result from interannual variations in temperature and precipitation (i.e., weather), which may be in part the result of long-term climate change, but there is yet no evidence for large permanent increases in CH₄ emissions. For 2009, our preliminary estimate of the global increase is ~4 ppb.

When evaluating these recent changes in atmospheric CH₄ burden, it is useful to consider the long-term behavior of CH₄. About a decade ago, we showed that if OH concentrations had been constant over the period of our observations, then total global CH₄ emissions had been constant. The observed decrease in CH₄ growth rate is consistent with a system approaching steady state, where the time constant of that approach is the CH₄ atmospheric lifetime (τ). Is this model of long-term behavior still consistent with the data today? The approach to steady state for this system, with zeroth-order source and pseudo-first-order loss, is given by: $[\text{CH}_4](t) = [\text{CH}_4]_{ss} - ([\text{CH}_4]_{ss} - [\text{CH}_4]_0)e^{-t/\tau}$. By fitting this equation (green line) to the global averages, we find $[\text{CH}_4]_{ss} \approx 1799$ ppb and $\tau \approx 9$ yr. The lifetime determined from this fit is in good agreement with values determined independently based on methyl chloroform observations. For more than two decades, atmospheric CH₄ has looked like a system approaching steady state; super-imposed on the decline in growth rate are significant variations. Understanding the processes driving these variations will improve our ability to predict how CH₄ emissions in the Arctic will respond to changing climate. So far, the evidence suggests recent increases in CH₄ are part of natural climate variability and not evidence for strong climate feedbacks in the Arctic.

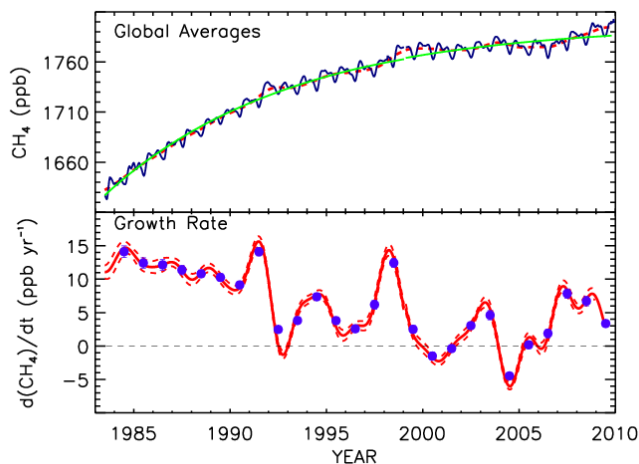


Figure 1. Globally averaged CH₄ mole fractions (blue), trend (red), and exponential fit (green; see text) (top panel); deseasonalized instantaneous growth rate (red) and annual increase (blue) (bottom panel). Results for 2009 are preliminary.