

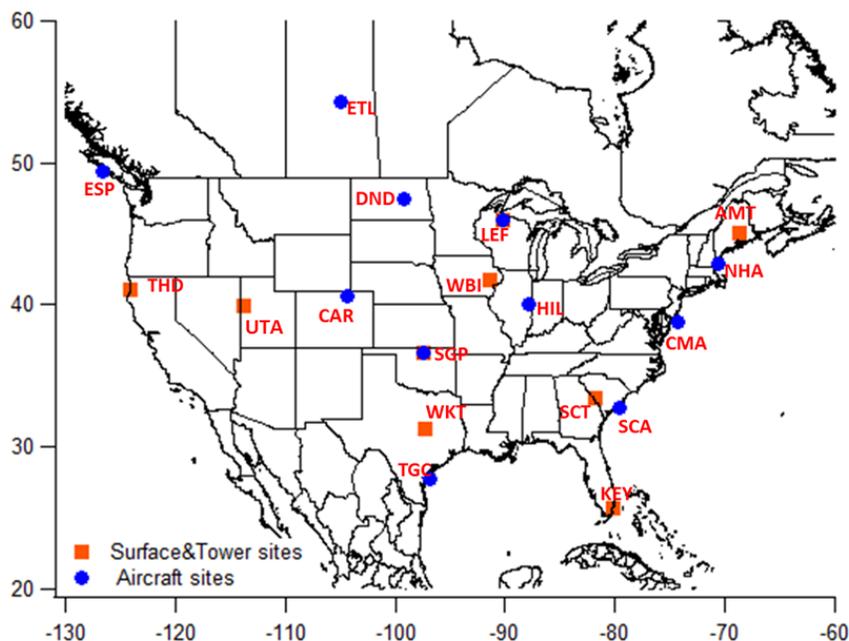
## Little Evidence for Significant Increases of CH<sub>4</sub> Emission in the U.S. over the Past Decade

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Recent studies on whether methane (CH<sub>4</sub>) emissions from oil and natural gas (ONG) operations in the U.S. have significantly increased are still inconclusive. To provide observational evidence we carefully analyzed the *in situ* CH<sub>4</sub> measurements from the NOAA Global Greenhouse Gas Reference Network (GGGRN) for the best estimates of CH<sub>4</sub> trends for 2006-2016. Methane data from 11 aircraft sites and 9 surface/tower sites across the U.S. were included in this study. Variations of sampling frequencies in different seasons were taken into account for accurate trend detection. We found that most of our sites had similar CH<sub>4</sub> trends of ~ 6.5 ppb/yr, which was comparable with the recent global background CH<sub>4</sub> trend from Mauna Loa. By utilizing the vertical gradient (Planetary Boundary Layer relative to high altitude data), we have the ability to detect significant increase of surface emissions larger than 7-10% over the study period from most of our sites. However, statistically significant increase were only found at the Southern Great Plain site in Oklahoma (SGP, downwind of the Eagle Ford, Barnett Shale and Woodford ONG fields) and the Dahlen sites in North Dakota (DND, downwind of the Bakken ONG field), which indicated influences from regional ONG activities. Ethane (C<sub>2</sub>H<sub>6</sub>) measurements from SGP (C<sub>2</sub>H<sub>6</sub> measurements were not available at DND) and propane (C<sub>3</sub>H<sub>8</sub>) measurements from both SGP and DND exhibited significant increasing trends. Linear correlations were well identified for surface C<sub>3</sub>H<sub>8</sub> and CH<sub>4</sub> enhancements at SGP, relative to observations at higher altitudes. However, by applying the observed enhancement ratios of surface C<sub>3</sub>H<sub>8</sub>/CH<sub>4</sub> and the C<sub>3</sub>H<sub>8</sub> trends (as indicator for ONG emissions) on CH<sub>4</sub> trend estimates, we would infer much larger surface CH<sub>4</sub> trends than what we actually observed at these two sites. When using C<sub>2</sub>H<sub>6</sub>, i-pentane, n-pentane, or n-butane data, we also infer much larger surface CH<sub>4</sub> trends. We found that the fat-tailed distribution of these hydrocarbon data and the changing enhancement ratios over time are likely responsible. This discrepancy suggests that using a constant enhancement ratio is not likely a reliable approach to compute CH<sub>4</sub> emission trends from ONG.



**Figure 1.** Long-term monitoring sites from the NOAA GGGRN.