







Radiative Surface Forcing from CH₄ at the North Slope of Alaska and Southern Great Plains Sites

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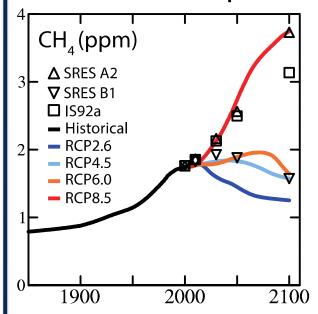
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- ⁴ Atmospheric and Environmental Research

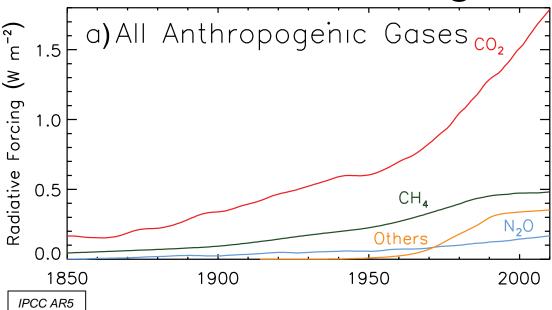






CH₄ Concentrations and Forcing





- While the concentrations are observed, these forcings are strictly modeled.
- Can we observe climate forcing?

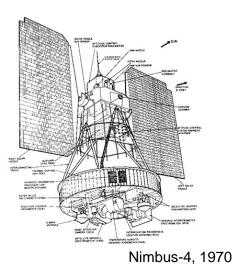




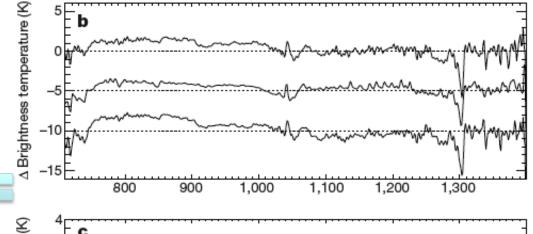
1st Detection of WMGHG Forcing

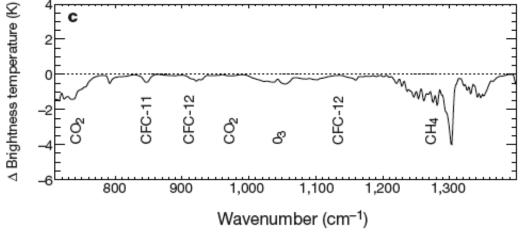


minus



Reduced Emission to Space from Enhanced Greenhouse

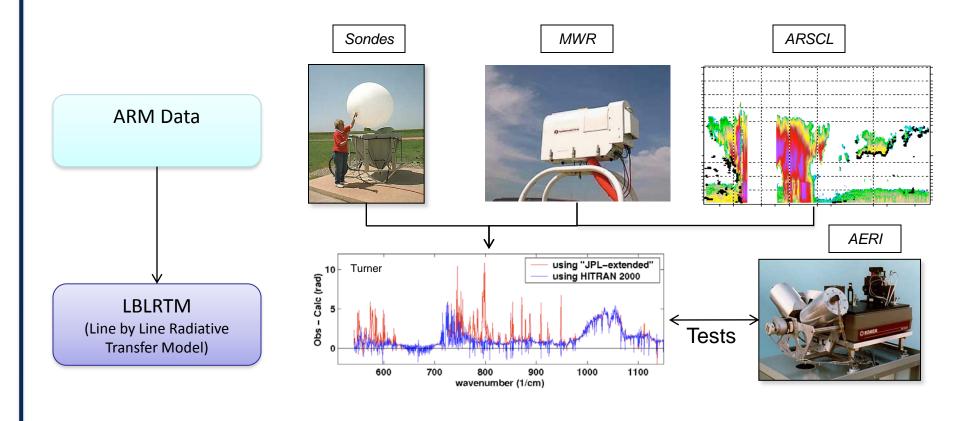








Schema for our Forcing Calculations



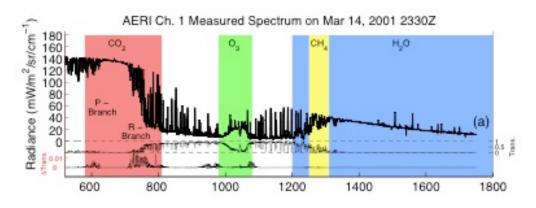
Feldman et al, Nature, 2015

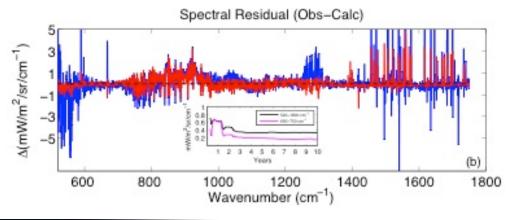


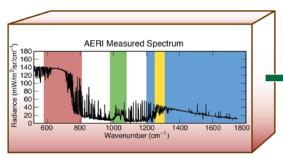


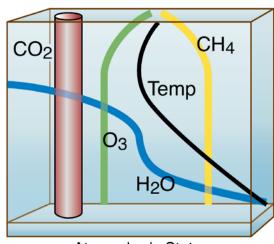
Measurements of CO₂ Surface Forcing

 ARM facilities at NSA and SGP contain numerous datastreams that enable the detection of the change in the surface energy budget due to rising CO₂.









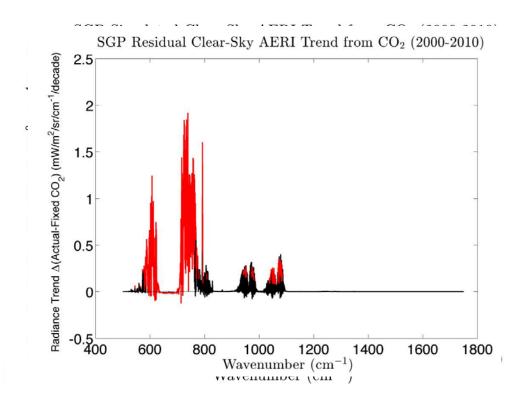
Atmospheric State

Feldman et al, Nature, 2015



Spectroscopic Separation

- Necessary to control for thermodynamic structure of the atmosphere through RAOBs.
- CO₂ absorption features are separable and appear in the measurements.



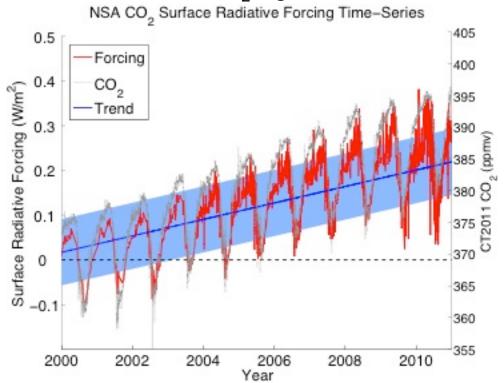
Feldman et al, Nature, 2015

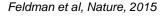




CO₂ Surface Forcing Time-Series

- We track systematic differences between AERI residuals (obs calc, fixed CO₂) between 2000 and 2010.
- Time-series at two sites reveal positive slopes ~0.2 W/m²/decade.
- Seasonal cycle ~ 0.5 W/m². Diurnal, synoptic, and seasonal features.
- Observational demonstration of CO₂'s greenhouse effect in action.



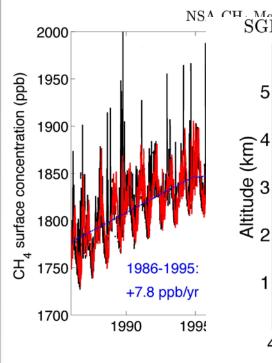


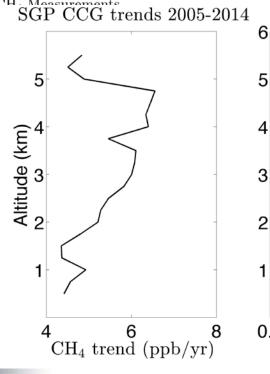


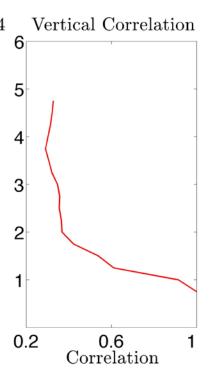


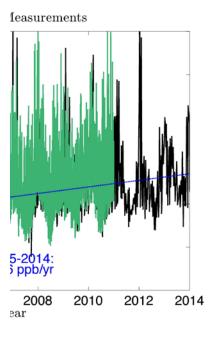
CH₄ Concentrations

- The hiatus in increasing atmospheric CH₄ ended ~2005.
 Measurements at NSA and SGP indicate this.
- Measured profiles of CH₄ also indicate trends in CH₄, but low vertical correlation.







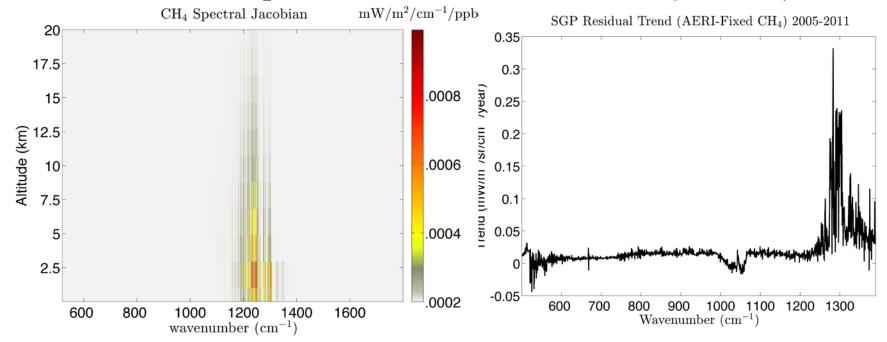






Trends in AERI Radiances – Fixed CH₄

CH₄ has a prominent absorption feature at 1311 cm⁻¹. Using the same approach as for CO₂, we can derive a time-series of forcing from CH₄.



Difference between AERI and model spectra with fixed CH₄ reveals:

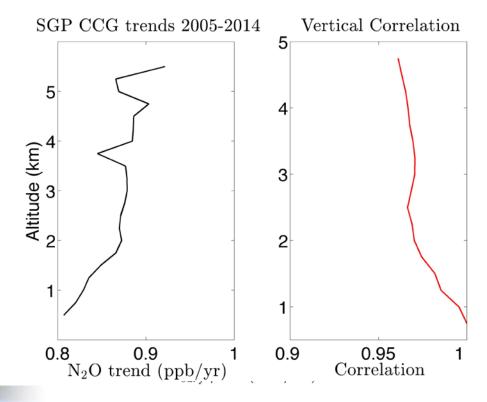
- Large secular trend in increased atmospheric emission in 1200-1350 cm⁻¹.
- Minor features outside of 1200-1350 cm⁻¹.





Challenges for CH₄ Forcing

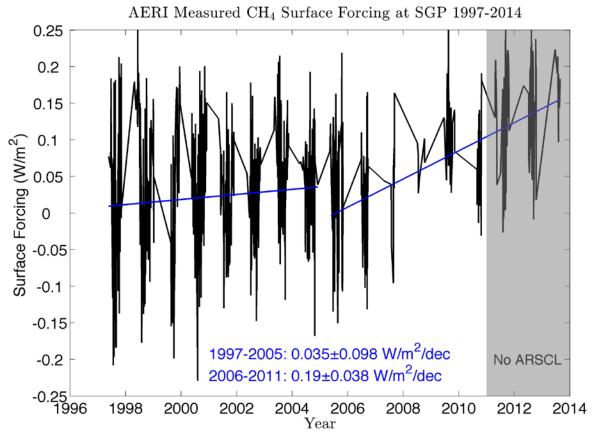
- CH₄ band is very sensitive to T and H₂O.
- N₂O and CH₄ features overlap.
 - But observations indicate N₂O uncorrelated with, and rising more slowly than, CH₄.







1st Detection of CH₄ Surface Forcing



Observational Determination of:

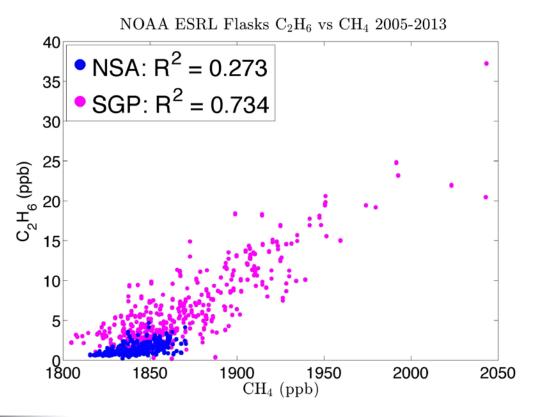
- •Time series of radiative forcing by CH₄: Signals of "hiatus" and recent rise.
- •Quantification of impact of CH₄ on the Earth's surface energy budget.
- •Determination of the radiative signal from seasonal cycle.





Anthropogenic Forcing

 Coincidental measurements of C₂H₆ and CH₄ indicate that spikes in surface CH₄ are generally due to anthropogenic activities, likely from a nearby source.







Conclusions

- The ARM data record is sufficiently long to quantify spectral signatures of anthropogenic forcing.
- We have used this record to quantify trends in surface radiation from recent changes in methane concentrations, with high-frequency spikes likely from anthropogenic activity.
- Trends: 0.19 ± 0.04 Wm⁻² / decade at the Southern Great Plains site.

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