Estimating Methane Emissions Consistent with Both Satellite and Isotope Constraints

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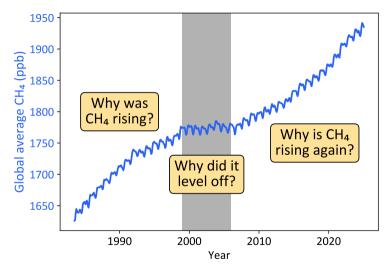




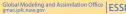


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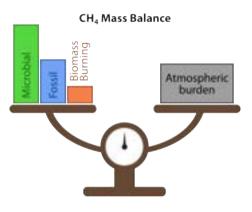


The leveling off of CH₄ in the early 2000's and its renewed growth after 2007 is a mystery with many possible explanations, but finding the right one is important for understanding current and future methane emissions and mitigation potential





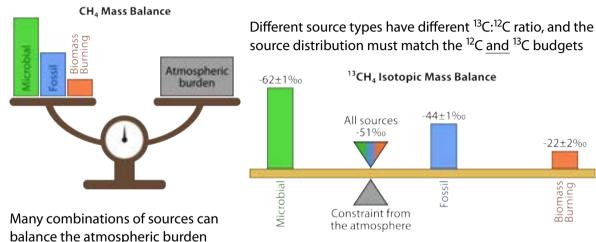




Many combinations of sources can balance the atmospheric burden





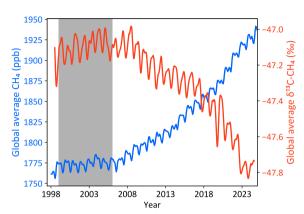


We have constructed an atmospheric inverse model with TM5 4DVAR to assimilate CH₄ and δ^{13} CH₄ measurements and estimate source-specific emissions





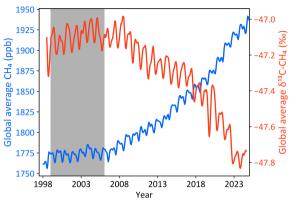




Through most of the 20^{th} century, $\delta^{13}CH_4$ has been increasing due to increasing fossil emissions. After ~2007, the trend reversed, pointing to influence of lighter sources.

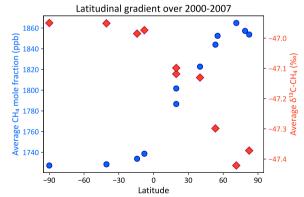






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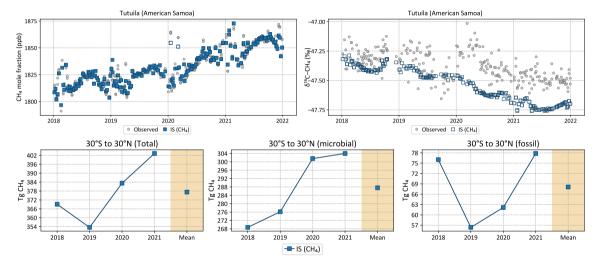
 $\delta^{13}\text{CH}_4$ is lower in the Northern Hemisphere because the average $\delta^{13}\text{CH}_4$ of all sources is lower than the atmosphere (CH₄ oxidation makes it heavier)





A CH₄-only inversion yields incorrect source partitioning



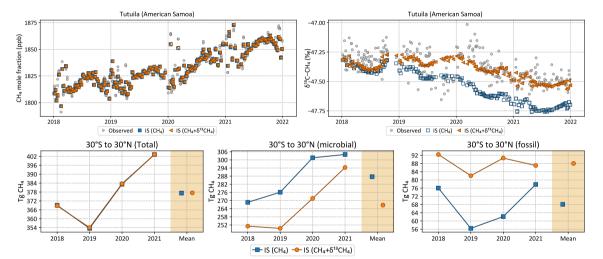


A CH₄-only inversion is not guaranteed to fit atmospheric δ^{13} CH₄ measurements, so unlikely to have the right partitioning between different source types unless the priors are already correct



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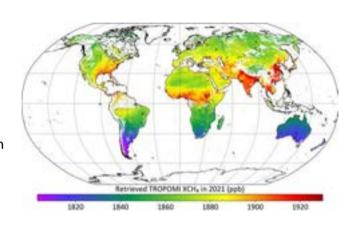
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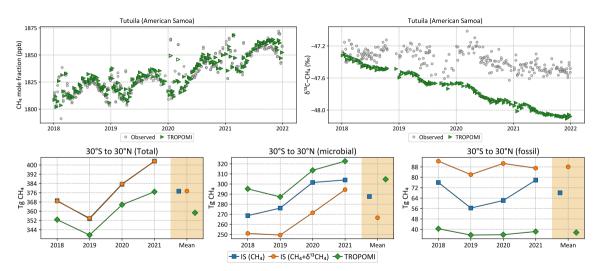


- SWIR XCH₄ retrievals starting 2018
- Current data selection:
 - Use only land data
 - Assimilate all retrievals with OA >= 0.5
 - Exclude snow & ice land retrievals. (Lindavist et al, 2024)
- High data density (\sim 300,000/day) leads to highly correlated retrieval errors, so inflate XCH₄ error based on the number of neighbors within 100 km and 30 min
- » Isn't source-specific except when different source types are spatially separated



Are TROPOMI retrievals consistent with in situ CH₄ and δ^{13} CH₄ obs?

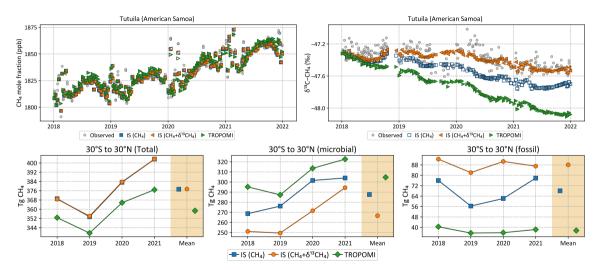




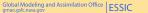


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TROPOMI and $\delta^{13}CH_4$ pull the source mixture in opposite directions, at least in the Tropics

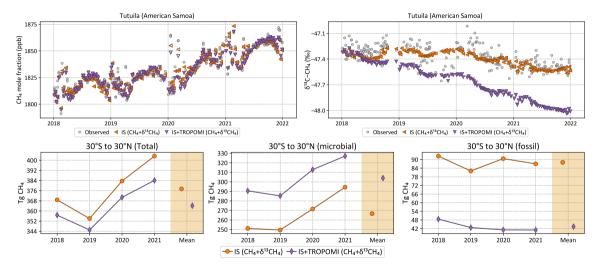




In a 1-box model of the atmosphere, this would be no problem. TROPOMI would provide a very strong constraint on the total CH₄ emissions, while δ^{13} CH₄ would provide information to split that into different source types.

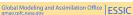
What happens when we try with the real atmosphere?





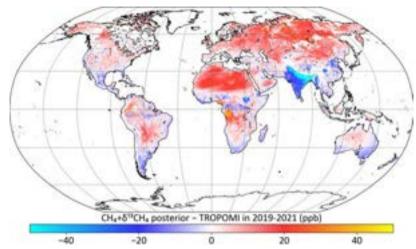
The partitioning is "wrong enough" with TROPOMI data that in situ $\delta^{13}CH_4$ can't fix it







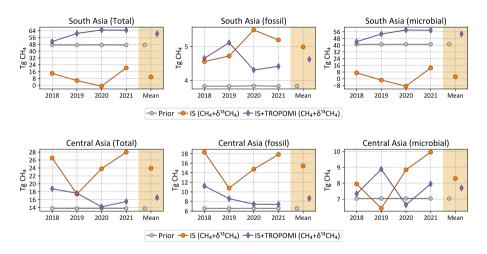




Adding TROPOMI XCH₄ will increase emissions over the blue regions and decrease them over the red regions, compared to an insitu-only CH₄+ δ^{13} CH₄ inversion







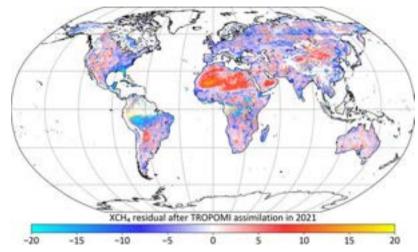
The total CH_4 estimates from TROPOMI are probably more accurate, but the source-specific adjustments are restricted by the priors







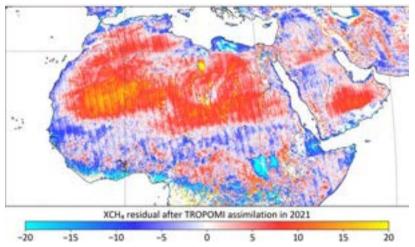




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TROPOMI XCH₄ seems to have a low bias over high-albedo high-aerosol scenes, and a weird striping pattern that cannot exist in column CH₄







- » Mathematically, the join assimilation of TROPOMI XCH₄ and in situ δ^{13} CH₄ is feasible. In theory, XCH₄ should provide a tight constraint on total CH₄ emissions, while δ^{13} CH₄ should split it among source types.
- » In practice, TROPOMI assimilation is consistent with in situ CH $_4$ but not δ^{13} CH $_4$
- » We should be checking satellite-derived CH₄ fluxes, especially their sectoral attributions, against data such as δ^{13} CH₄ that provide independent sectoral information



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- » We should be checking satellite-derived CH₄ fluxes, especially their sectoral attributions, against data such as δ^{13} CH₄ that provide independent sectoral information
- » Part of the problem is that the relative magnitude of priors in some parts of the world are not consistent with atmospheric $\delta^{13}\text{CH}_4$
- $^{>\!>}$ TROPOMI XCH $_4$ retrievals also seem to have albedo dependent artifacts and some sort of across-orbit bias pattern (striping), working on better data selection



