

Towards a remote sensing solution to quantify N₂O emissions by integrating shortwave and longwave infrared bands

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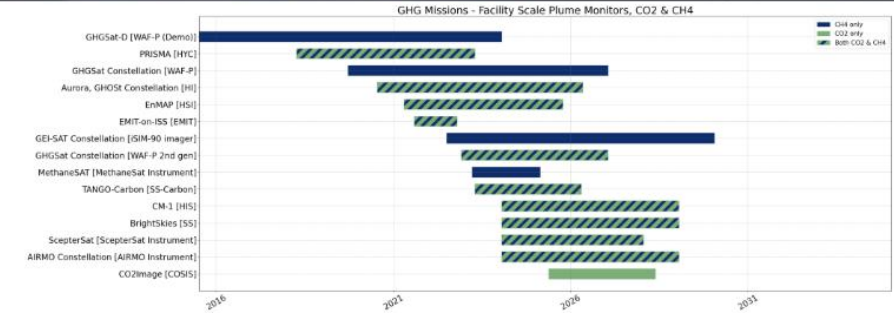
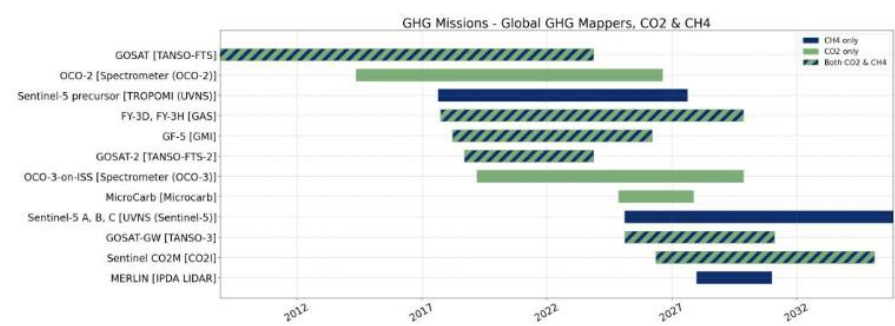
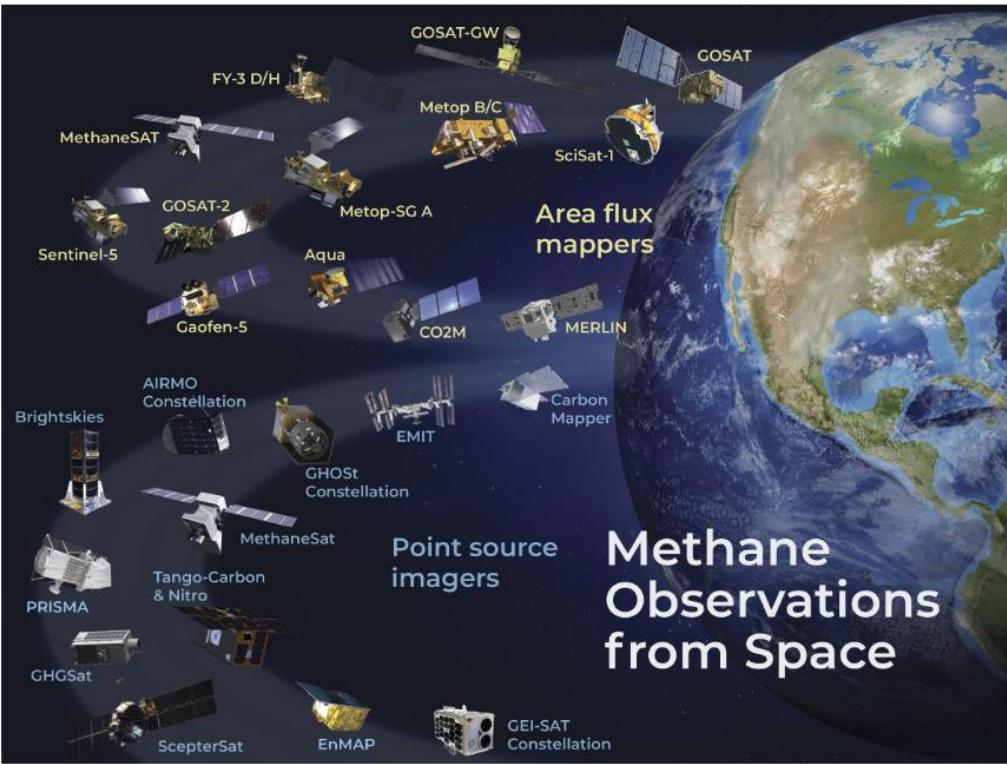
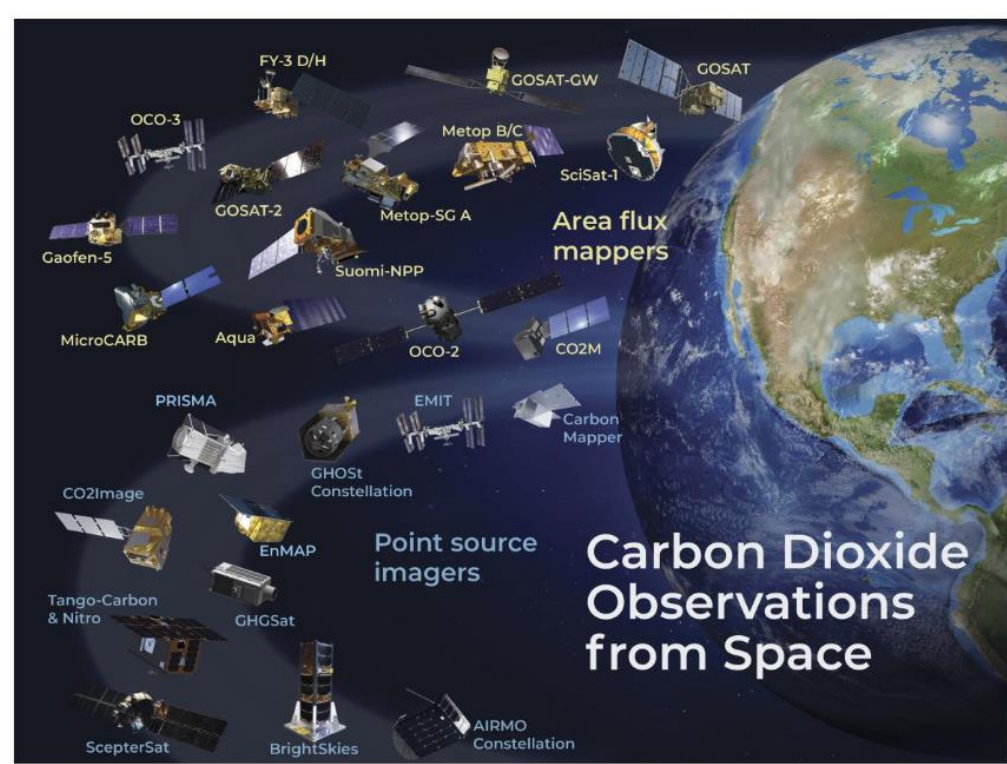
Radiative transfer modelling: Christopher Chan Miller, Robert Spurr, Karen Cady-Pereira

Instrument design: Thomas U. Kampe, Nathan Leisso

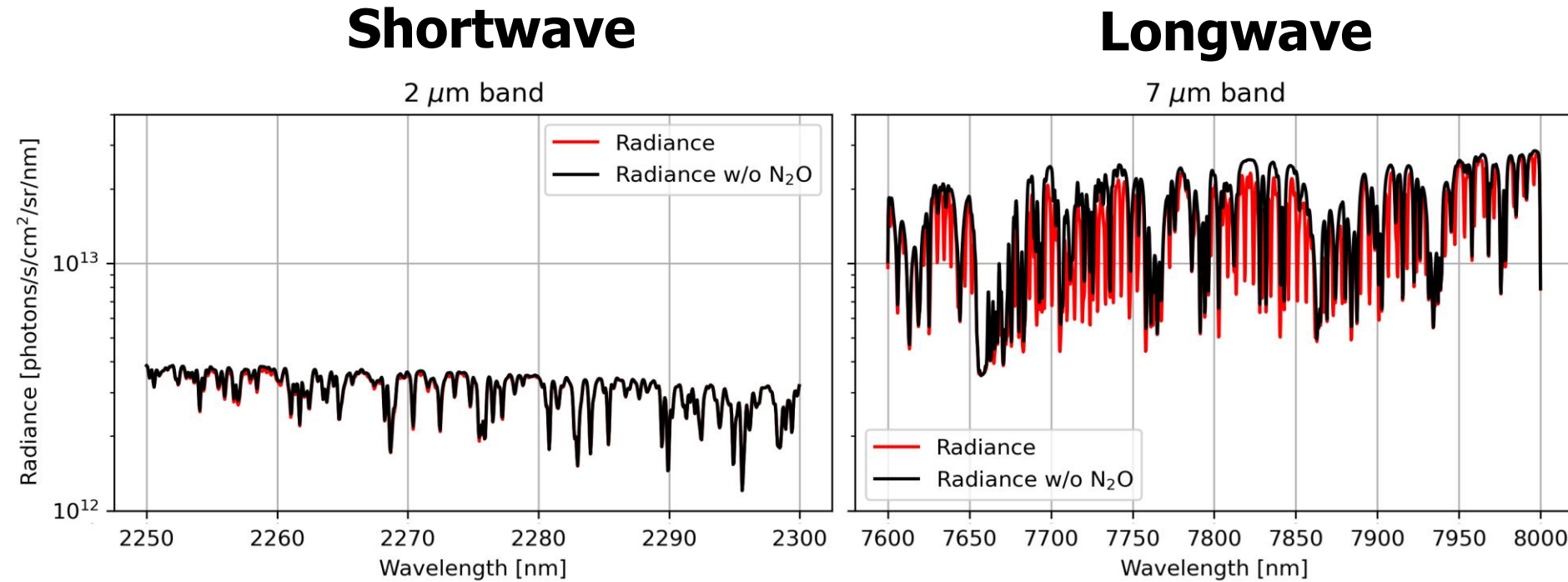
N₂O flux data: Wendy H. Yang, Emily R. Stuchiner, Eddy Will



N₂O: a key greenhouse gas overlooked from space



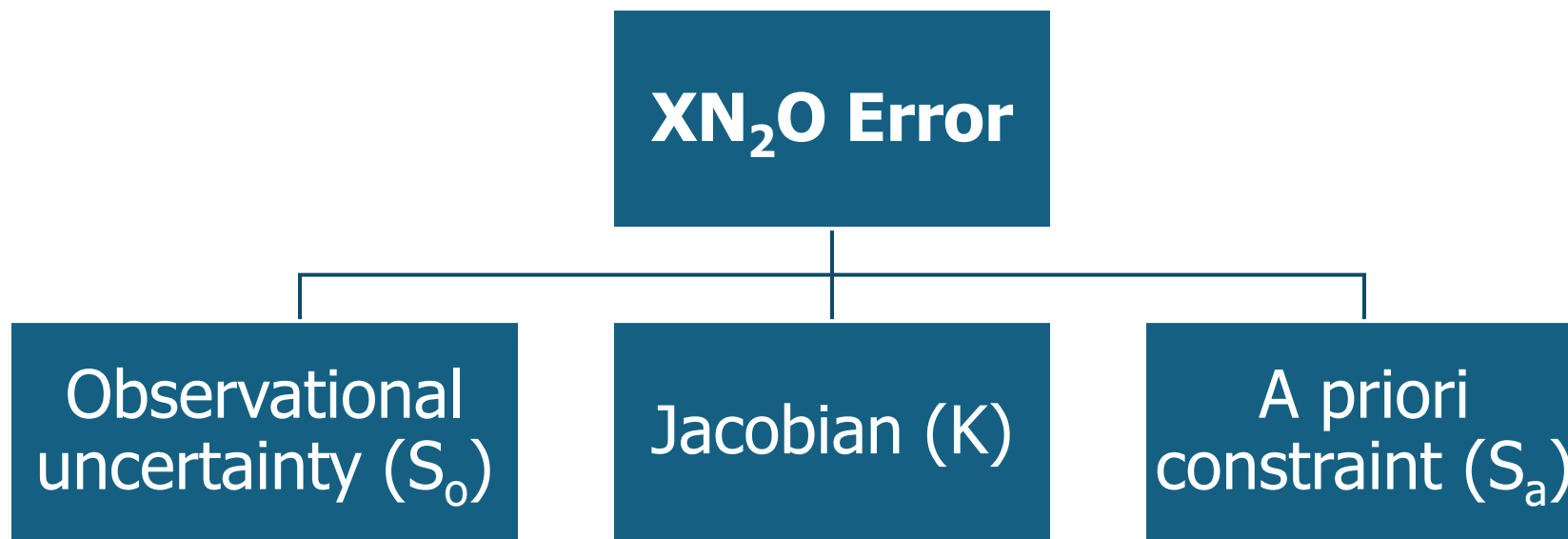
The challenge of N₂O remote sensing



- 2 μm : consistent vertical sensitivity but very weak N₂O absorption
- 7 μm : strong absorption and radiance level but weak near-surface sensitivity
- Integrate 2 μm and 7 μm to combine the strengths of short and longwave bands

How precisely can we observe XN_2O ?

- XN_2O : Column-integrated mixing ratio of N_2O $(\text{XN}_2\text{O} = \frac{\Omega_{\text{N}_2\text{O}}}{\Omega_{\text{A}}})$



\mathbf{S}_m : measurement error covariance matrix

Measurement error of XN_2O

$$\mathbf{S}_m = \mathbf{G}\mathbf{S}_o\mathbf{G}^T$$

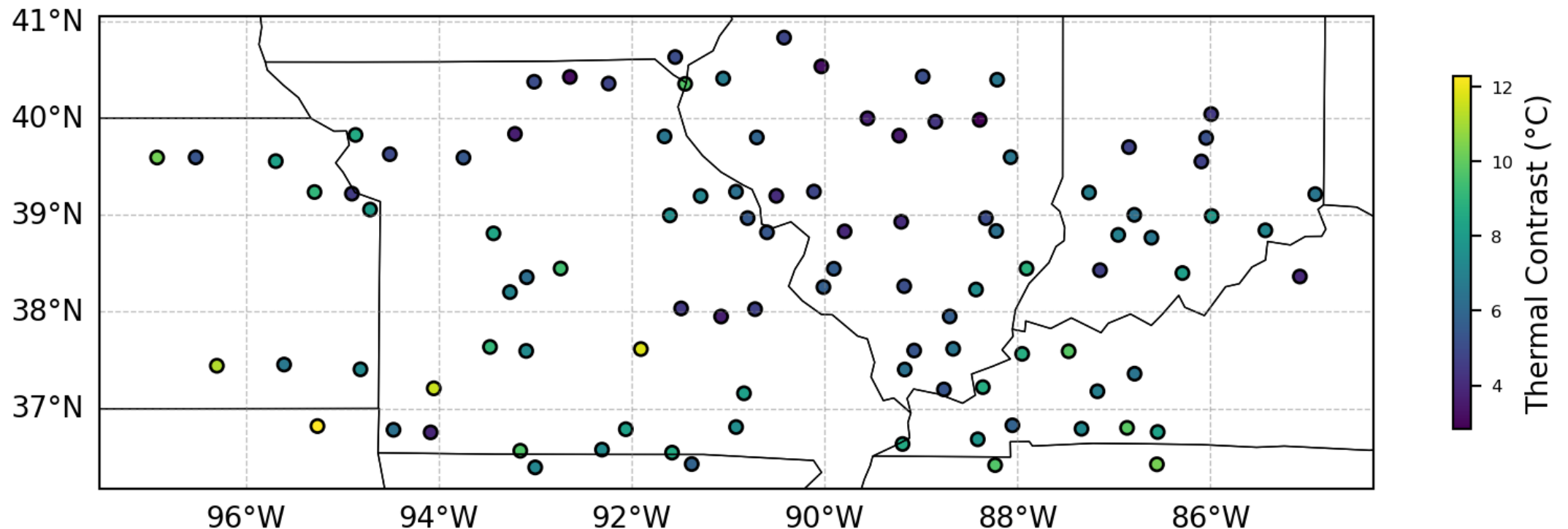
$$\sigma_{\text{XN}_2\text{O}(m)} = \mathbf{h}^T \mathbf{S}_m \mathbf{h}$$

$$\mathbf{G} = \mathbf{S}_a \mathbf{K}^T (\mathbf{K} \mathbf{S}_a \mathbf{K}^T + \mathbf{S}_o)^{-1}$$

\mathbf{h} : pressure weighing function

Atmospheric and surface states

- CrIS Level 2 files for N_2O , CH_4 , H_2O , and temperature from the JPL MUSES algorithm
- Atmospheric profiles and surface conditions extracted from 100 CrIS pixels
- Area: US Midwest (Part of corn belt)
- Observation date: 23 August 2023



Observational uncertainty

Parameter [unit]	2 μm band	7 μm band
Wavelength range [nm]	2240–2300	7600–8000
Spectral sampling [nm]	0.0575	0.25
Slit width [× spectral sampling] ^a	3	3
Exposure time [s]	0.1	0.1
Detector pixel size [μm]	18	18
f-number	2	2
System efficiency	0.5	0.5
Readout noise [electrons]	60	60
Airborne observation altitude [km]	10	10
Spaceborne observation altitude [km]	600	600
Airborne along-track [m]	20	20
Spaceborne along-track [m]	701	701
Airborne across-track [m]	2.95	2.95
Spaceborne along track [m]	155	155
Airborne ground sampling distance [m]	7.7	7.7
Spaceborne ground sampling distance [m]	330	330

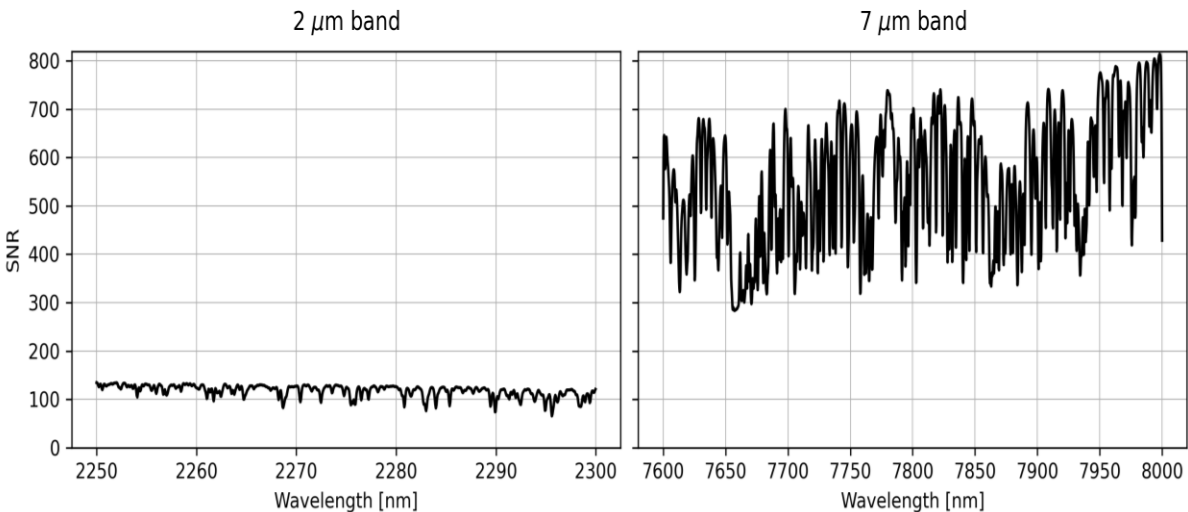
^a Gaussian slit function (i.e., instrument spectral response function, ISRF) is assumed.

$$S = \frac{\pi}{4} \left[I \right] dp^2 \cdot \frac{1}{f^2} \cdot n_{\text{sample}} \cdot dt \cdot d\lambda \cdot \eta$$

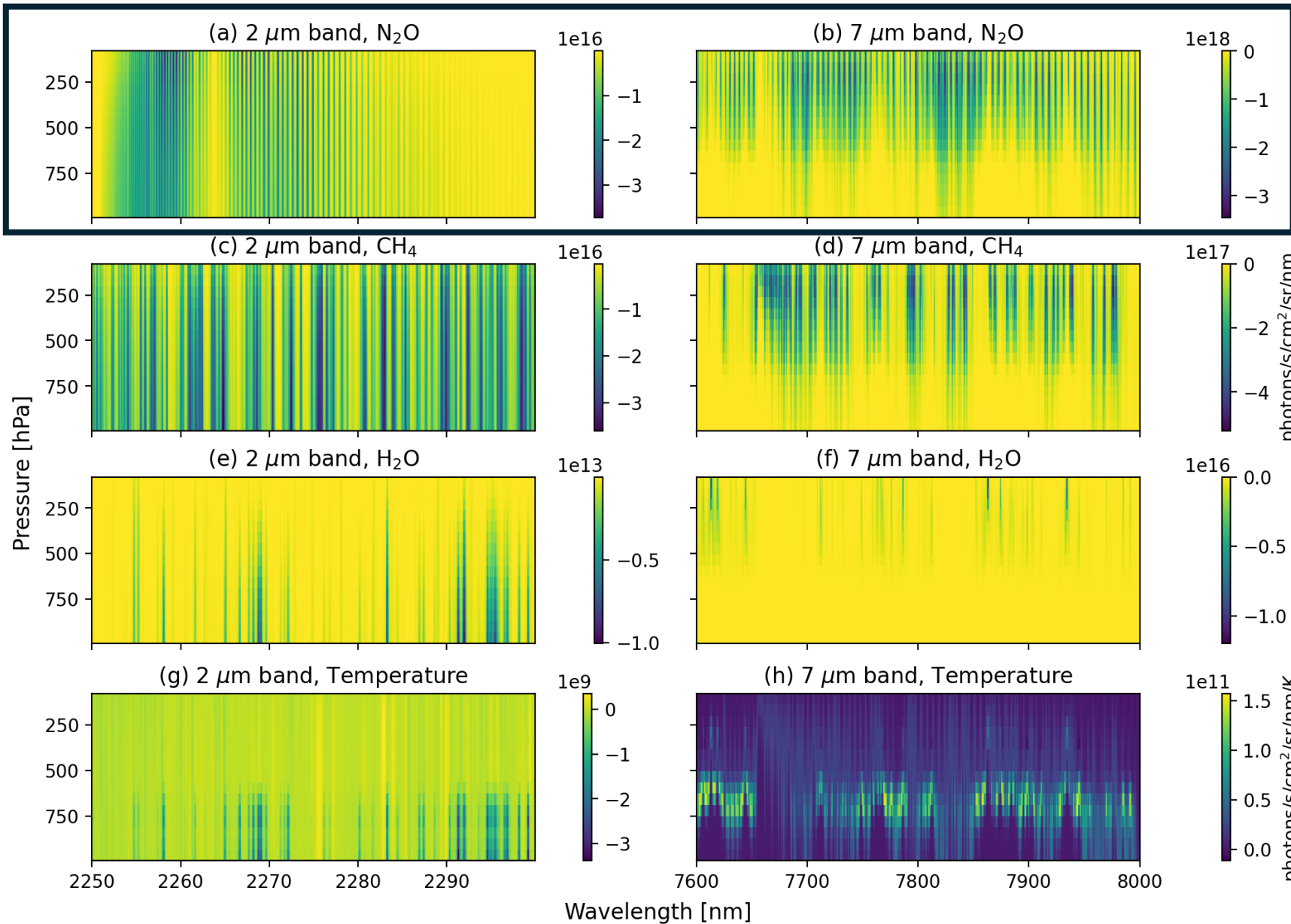
$$N = \sqrt{S + N_r^2}$$

$$\text{SNR} = \frac{S}{N}$$

SPLAT-VLIDORT radiative transfer model is used to simulate radiance (I)

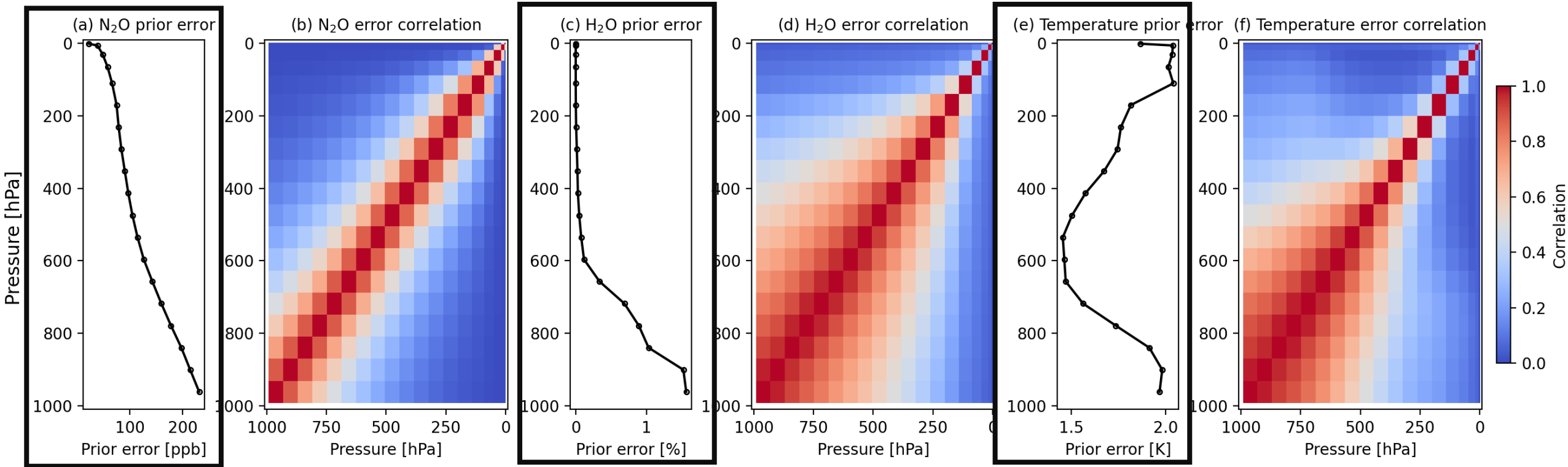


Radiance sensitivity to state vector (Jacobians)



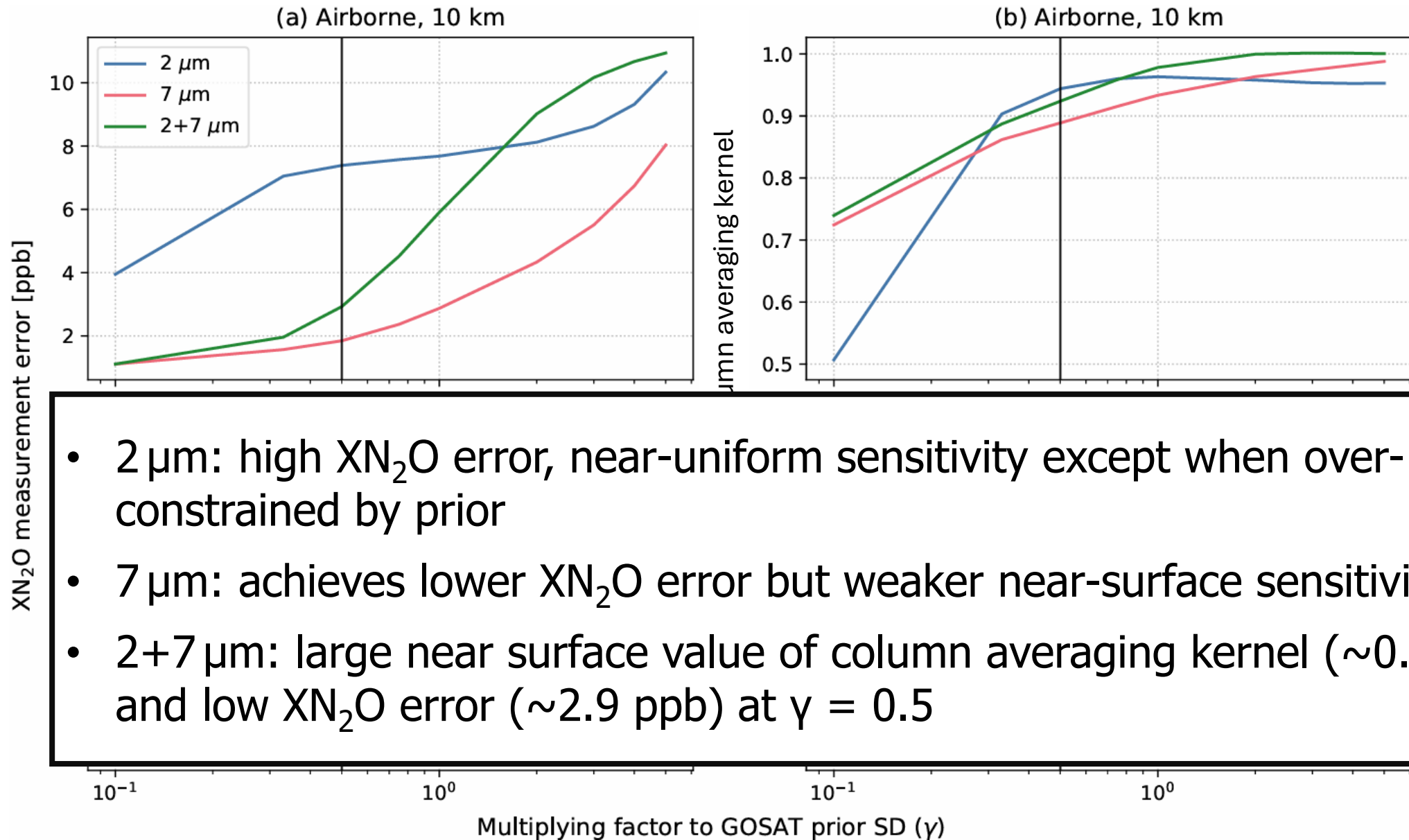
- State vector includes N_2O , CH_4 , water vapor, temperature profiles, and surface temperature
- 2 μm : sensitivity to the whole column including near-surface
- 7 μm : stronger sensitivity to N_2O as compared to the 2 μm band, but less sensitive to near-surface layers

A priori constraint



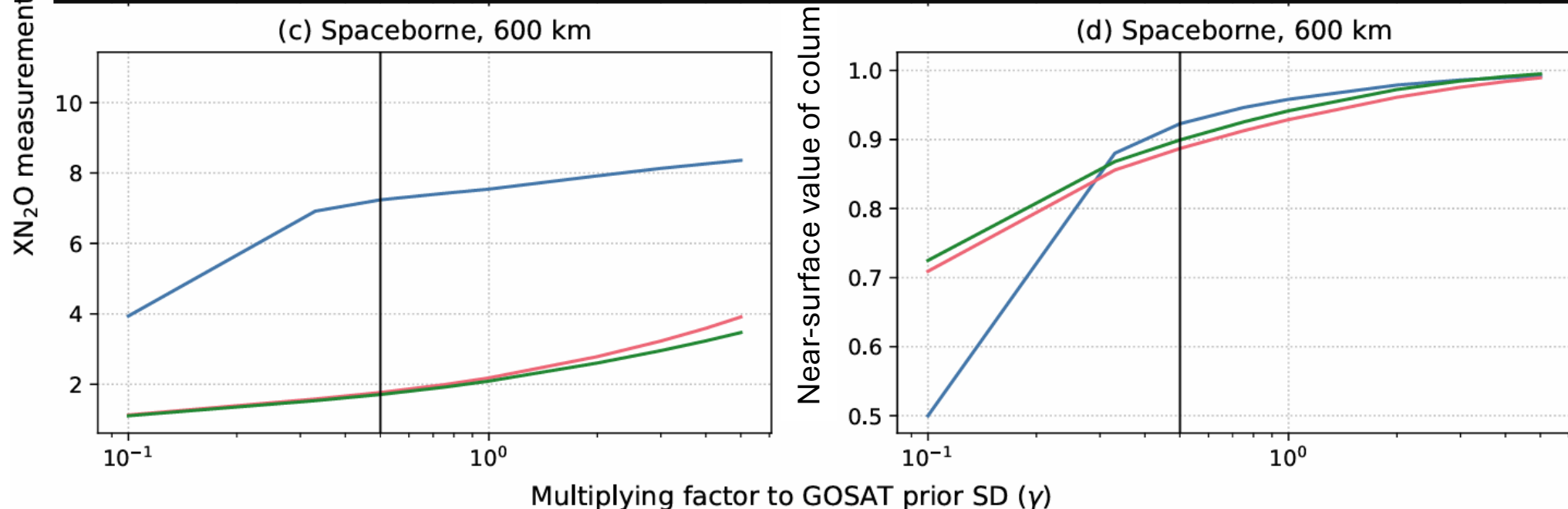
- N₂O prior error is derived from University of Leicester GOSAT CH₄ retrieval, scaled by their background concentration ratio (330 ppb / 1900 ppb)
- H₂O and temperature priors are adopted from CrIS algorithm
- Scaling factor gamma (γ) is applied to N₂O prior standard deviation to tune prior constraint strength

XN₂O precision



XN₂O precision

- 2 μm : highest XN₂O error across all prior strengths
- 7 μm : comparable XN₂O error with dual-band approach but weaker near-surface sensitivity
- 2+7 μm : large near surface value of column averaging kernel (~ 0.9) and low XN₂O error (~ 1.7 ppb) at $\gamma = 0.5$



How precisely can we observe XN_2O ?

Using 2 μm and 7 μm integration and at a moderate prior strength ($\gamma = 0.5$), we achieve:

- 2.9 ppb XN_2O error for airborne
- 1.7 ppb XN_2O error for spaceborne
- ~ 0.9 near surface value of column averaging kernel for both

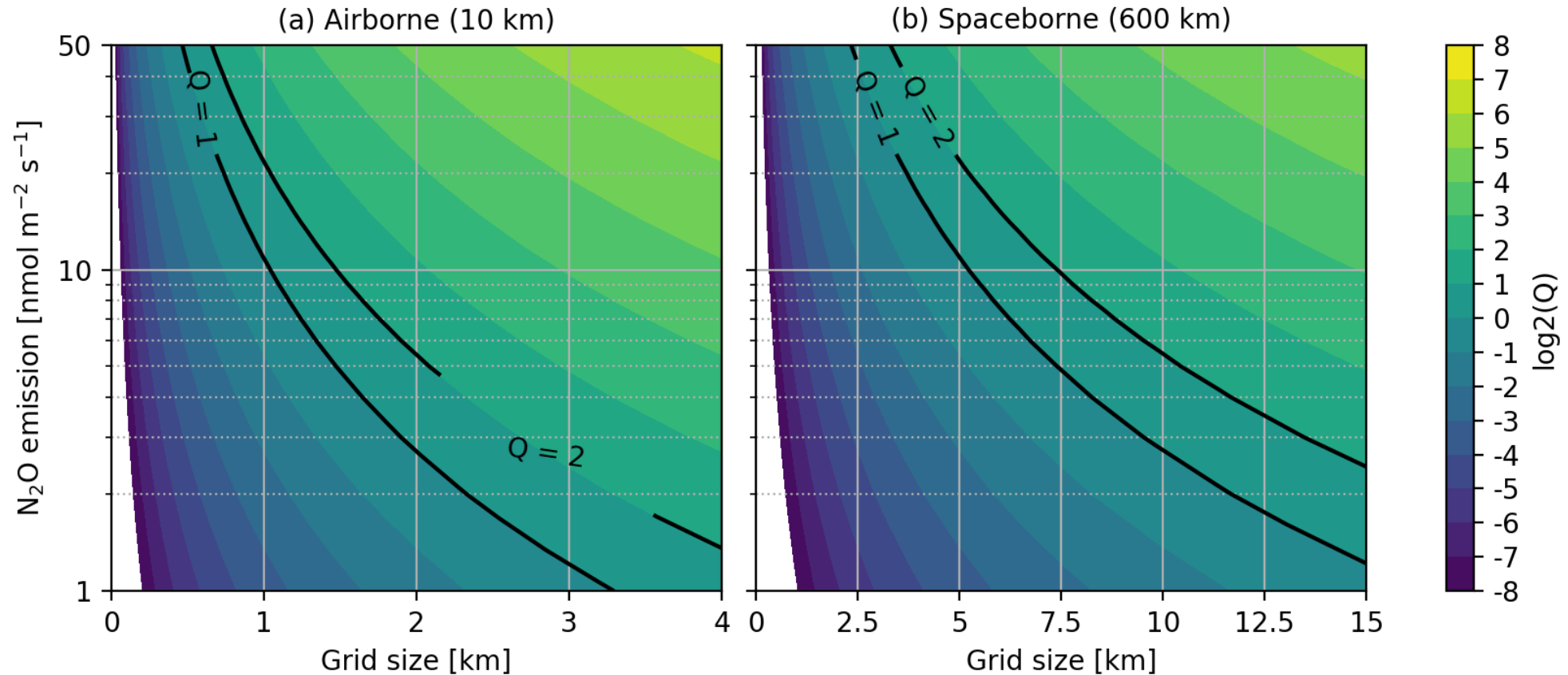
With those precisions, can we detect actual surface emissions, and at what spatial scale?

Estimating N₂O emissions using errors

$$Q = \frac{\text{Emission induced enhancement } (\Delta\Omega)}{\text{Column error } (\sigma_\Omega)} = \frac{E\Delta x^2 M_{\text{air}} g}{|u| \sigma_{\text{XN}_2\text{O}} x_0 P_{\text{air}}}$$

Parameter	Value [unit]
Emissions (E)	1-50 [nmol m ⁻² s ⁻¹]
Airborne ground sampling distance (x _{0_air})	7.7 [m]
Spaceborne ground sampling distance (x _{0_space})	330 [m]
Airborne aggregation scale (Δx)	X _{0_air} to 4 [km]
Spaceborne aggregation scale (Δx)	X _{0_space} to 15 [km]
Molar mass of dry air (M _{air})	0.029 [kg mol ⁻¹]
Acceleration due to gravity (g)	9.8 [m s ⁻²]
Wind speed (u)	1.389 [m s ⁻¹]
Airborne σ _{XN₂O}	2.9 [ppb]
Spaceborne σ _{XN₂O}	1.7 [ppb]
Pressure (P _{air})	1e5 [Pa]

Estimating N₂O emissions using errors



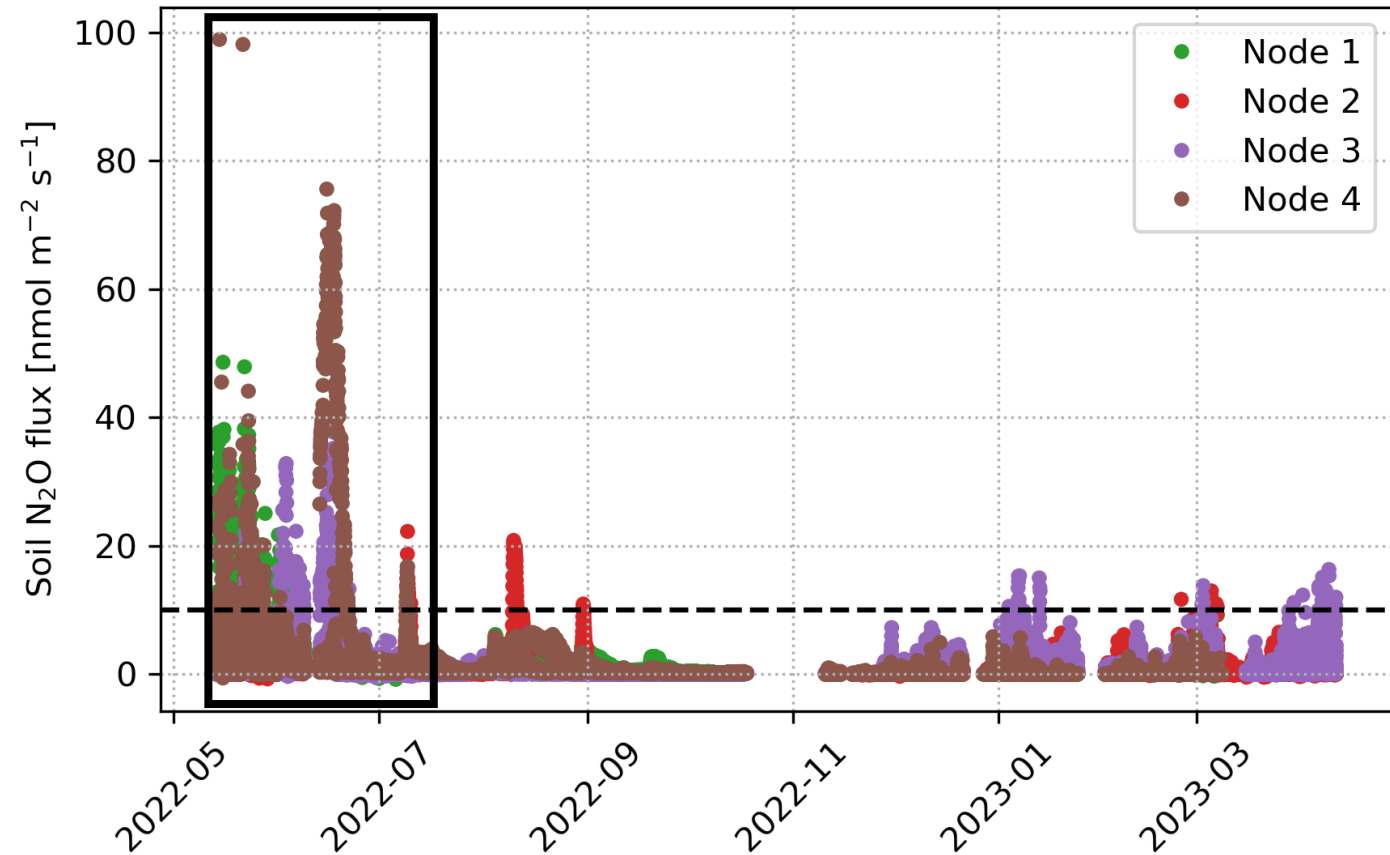
- Airborne: 10 nmol m⁻² s⁻¹ emission flux is observable at unit signal-to-noise ratio (Q) at ~1 km
- Spaceborne: 10 nmol m⁻² s⁻¹ emission flux is observable at unit signal-to-noise ratio (Q) at ~5 km

N₂O chamber data from University of Illinois



Auto-chambers on conventionally tilled maize field in central Illinois, USA

- 4 Nodes, 5 chambers per node
- $5 \times 10^4 \text{ m}^2$ area of the field, with nodes 50–100 m distance apart
- $10 \text{ nmol m}^{-2} \text{ s}^{-1}$ or higher N₂O emission does occur in the real field especially during growing season
- N₂O flux correlated for nodes that are 100 m apart



Conclusion

- Evaluated remote sensing solutions for high-resolution mapping of N₂O flux variability in agricultural landscapes
- Integrating the shortwave and longwave bands deliver strong near-surface sensitivity ($AVK \approx 0.9$) and low XN₂O error (≈ 2.9 ppb airborne; 1.7 ppb spaceborne)
- Emission flux of $10 \text{ nmol m}^{-2} \text{ s}^{-1}$ is observable at $Q = 1$ down to ~ 1 km for airborne and ~ 5 km for spaceborne