



Long-term trends and spatial distributions of CO₂, CH₄, N₂O and SF₆ from NOAA's global background measurements

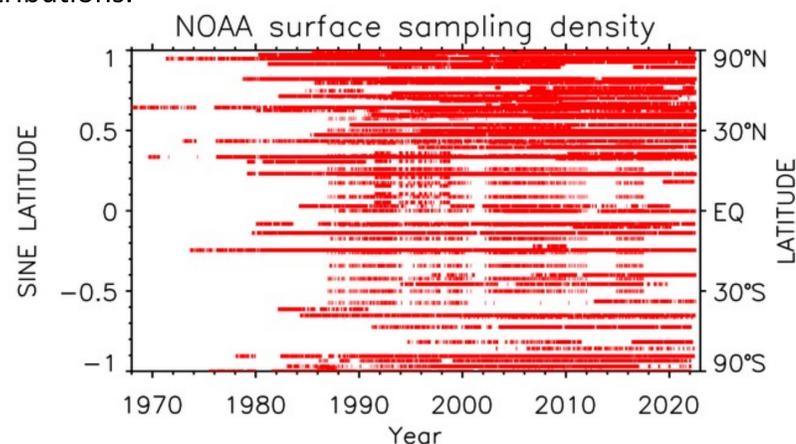


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Introduction

NOAA's Global Monitoring Laboratory began monitoring CO₂ from weekly discrete air samples collected at Niwot Ridge, Colorado and Ocean Station M in 1968. Since then, the network of flask-air sampling sites has grown, with more than 50 active sites in early-2022, and we now precisely measure CO₂, CH₄, N₂O, and SF₆. The goal of this network is to track the GHG abundances, determine budgets and understand carbon cycle feedback. Success or failure at meeting this goal depends the long-term continuity of measurements with sufficient network sampling density, but most importantly, it depends on the measurement quality as whether they can provide precisely measure on spatial and temporal distributions.



Analytical Capabilities (flask-air)

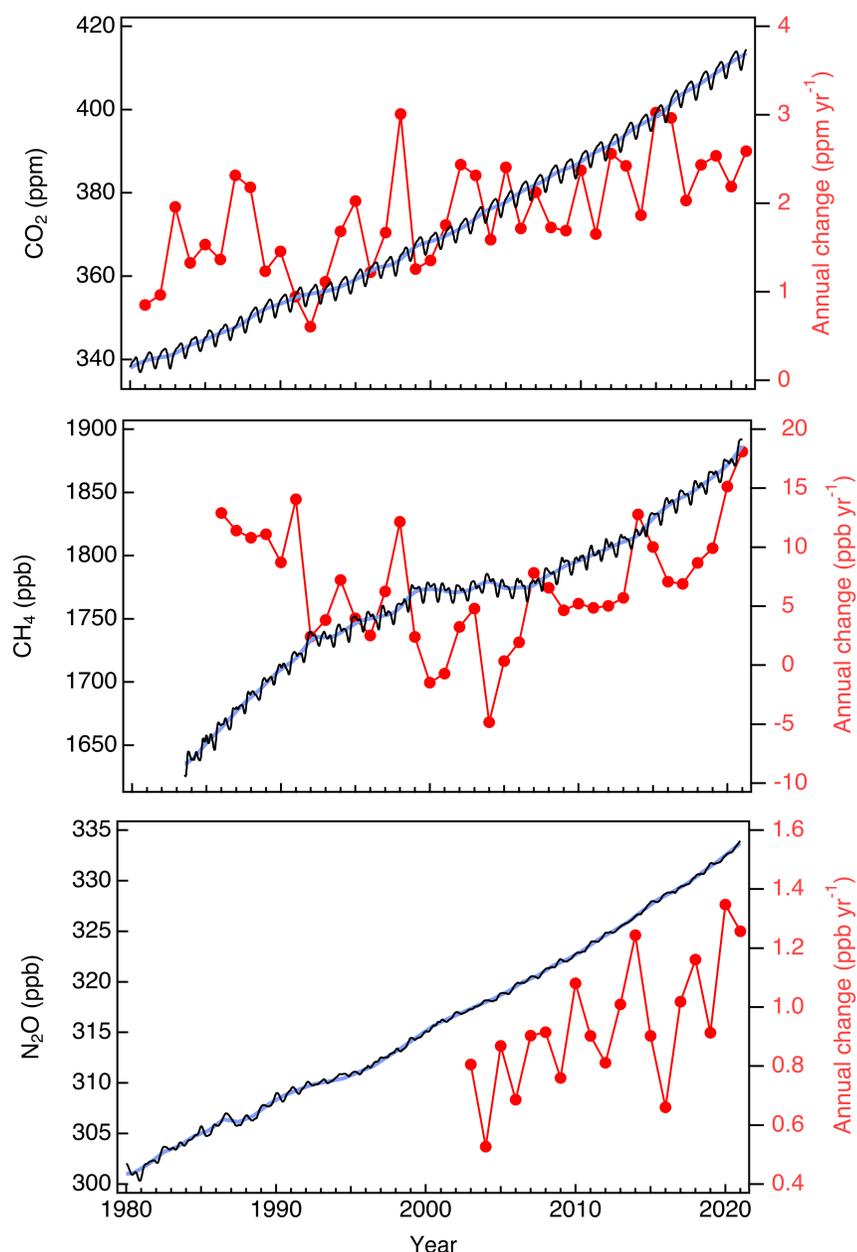
Our network delivers internally-consistent, calibrated observations over long time scale by enforcing detailed QA/QC procedures

Gas	Uncertainty (68% CI)	Technique
CO ₂	0.07 → 0.04 μmol mol ⁻¹ (ppm)	NDIR → CRDS
CH ₄	0.9 → 0.6 nmol mol ⁻¹ (ppb)	GC/FID → CRDS
N ₂ O	0.26 → 0.16 nmol mol ⁻¹ (ppb)	GC/ECD → TILDAS
SF ₆	0.04 pmol mol ⁻¹ (ppt)	GC/ECD
δ ¹³ CO ₂	*0.01‰	DI-IRMS
δ ¹³ CH ₄	*0.06‰	GC/CF-IRMS

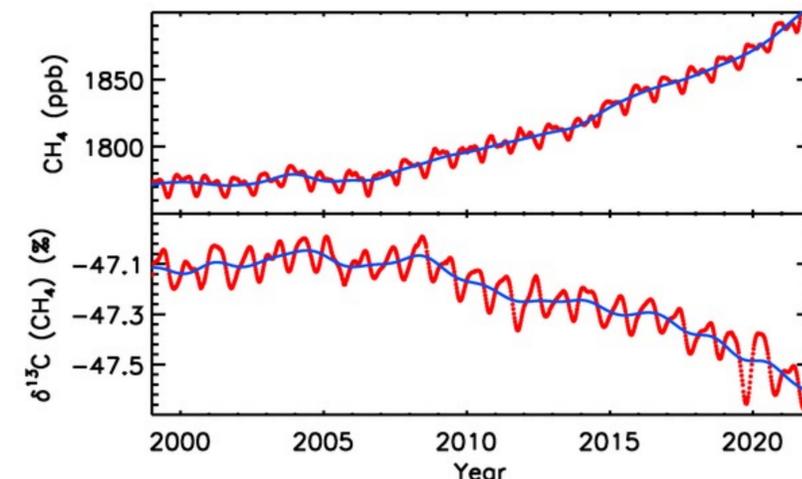
Uncertainty includes short-term repeatability, long-term reproducibility and scale propagation uncertainty

* Repeatability only

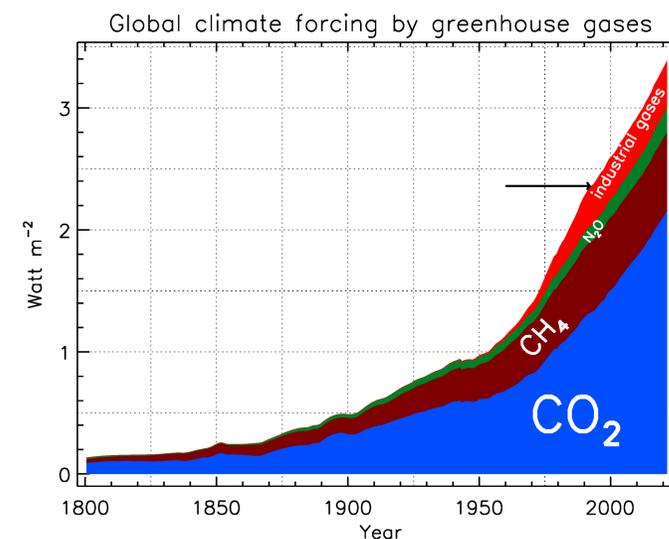
Atmospheric abundance



Global mean dry-air surface mole fractions (approximately weekly data in black, left axis) and annual change (red, right axis) derived from the NOAA Global Greenhouse Gases Reference Network. Deseasonalized trend curves are shown in blue. N₂O data prior to 2001 are too sparse to allow robust estimates of annual growth rates. Atmospheric abundances of these greenhouse gases have been increasing significantly in the past few decades



Measurements of CO₂ and CH₄ isotopes are available from our network to improve our understanding on the underlying emission/sink processes. The increasing trend of CH₄ coincide with the decreasing trend in δ¹³C-CH₄ indicates a dominant role of increasing microbial emissions after 2006.



Increase in atmospheric CO₂ contributes 66% of the increase in all GHGs radiative forcing since the pre-industrial era while the increase in CH₄ contributes 16%. Over the last five years, the increase of instantaneous climate forcing by CO₂ has accounted for about 80% of the rate of increase of all GHG forcing. The increase of CH₄ has been a quatre as large as the increase caused by CO₂. The increase of forcing by N₂O over has been 70% as large as the increase caused by CH₄. Reduction of CO₂ emissions must remain the focus in mitigating climate change.