

**Improving
measurements of SF₆
for the study**

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Improving measurements of SF₆ for the study of atmospheric transport and emissions

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Abstract

Sulfur hexafluoride (SF_6) is a potent greenhouse gas and useful atmospheric tracer. Measurements of SF_6 on global and regional scales are necessary to estimate emissions and to verify or examine the performance of atmospheric transport models. Typical precision for common gas chromatographic methods with electron capture detection (GC-ECD) is 1–2%. A method for improving measurement precision is described. We have modified a common GC-ECD method to achieve measurement precision of 0.5% or better. Global mean SF_6 measurements were used to examine changes in the growth rate of SF_6 and corresponding SF_6 emissions. Global emissions and mixing ratios from 2000–2008 are consistent with recently published work. More recent observations show a 10% decline in SF_6 emissions in 2008–2009, which seems to coincide with a decrease in world economic output. This decline was short-lived, as the global SF_6 growth rate has recently increased to near its 2007–2008 maximum value of $0.30 \pm 0.03 \text{ pmol mol}^{-1} (\text{ppt}) \text{ yr}^{-1}$ (95% C.L.).

1 Introduction

Sulfur hexafluoride (SF_6) is a potent greenhouse gas with a global warming potential (100 yr time horizon) 22800 times that of carbon dioxide (Forster et al., 2007). Because of its very long lifetime (800–3200 yr) (Ravishankara, et al., 1993; Morris et al., 1995) and the fact that the only known sink is destruction in the mesosphere, the atmospheric burden is essentially equal to the total amount of SF_6 emitted during the industrial era. While some voluntary efforts to reduce the amount of SF_6 emitted to the atmosphere have been made (Harnisch and Gluckman, 2001; US EPA, 2008;) the atmospheric concentration continues to rise (Elkins and Dutton, 2009; Levin et al., 2010; Rigby et al., 2010). Efforts to quantify SF_6 emissions have been performed using bottom-up inventories and top-down methods based on atmospheric concentration and variability (UNFCCC, 2010; Maiss and Brenninkmeijer, 1998; Bakwin et al., 1997;

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Hurst et al., 2007; Levin et al., 2010; Rigby et al., 2010). Reconciling discrepancies between top-down and bottom-up emissions should, in principle, be relatively simple compared to other greenhouse gases for which there are natural sources or sinks that are more difficult to quantify (e.g. N₂O and CH₄). Still, discrepancies between bottom-up and top-down emissions of SF₆ remain (Levin et al., 2010), and estimating emissions on regional scales by top-down methods remains challenging (Rigby et al., 2010). Measurement precision is one of the limiting factors in estimating emissions through atmospheric inversion methods (Rigby et al., 2010). SF₆ is also used to evaluate atmospheric transport models (Levin and Hesshaimer, 1996; Gloor et al., 2007; Patra et al., 2009), and is particularly important for the study of stratospheric circulation and the calculation of air mass mean age (Park et al., 1999, Engel et al., 2008, Ray et al., 2010).

Measurements of atmospheric SF₆ are performed by a number of researchers around the world. The predominant method of SF₆ measurement involves gas chromatography with detection by electron capture detector (ECD) or mass selective detection (MSD). The ECD is sufficiently sensitive to SF₆ that pre-concentration is generally not required, although large volumes (5–25 cm³) are typically injected. Pre-concentration has been used with ECD detection (Wanninkhof, 1991) and also with mass selective detection (Miller et al., 2008). With the ECD, SF₆ is typically separated from other ECD-sensitive gases by packed columns, such as molecular sieve 5A (Simmonds et al., 1972), Porapak-Q, or Hayesep-Q (Elkins, 1980; Elkins et al., 1996; Hall et al., 2007; Thompson et al., 2009). Because nitrous oxide (N₂O) and SF₆ can both be resolved using Porapak-Q (or Hayesep-Q) columns, this method is commonly employed (Worthy et al., 2003; van der Laan et al., 2009; Thompson et al., 2009). Hayesep-Q and Porapak-Q are similar in their characteristics and result in similar chromatography. Typical precision for most ambient SF₆ measurements using ECDs is 1–2%. Better precision can sometimes be achieved using pre-concentration, but this adds additional complexity, and is not as common as loop-injection air samples.

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Here we present a modification of a common Porapak-Q method that combines Porapak-Q and molecular sieve 5A columns to achieve better SF₆ precision than is typically observed using Porapak-Q alone. We first describe the Porapak-Q system used to measure SF₆ in NOAA/ESRL flask samples of ambient air since 1995 (Geller et al., 1997). Then we describe how this method was modified to improve SF₆ precision on a laboratory-based GC used for calibration, and also on an in situ system. Finally, we use global and hemispheric mean SF₆ from NOAA/ESRL flask and in situ networks to examine recent changes in emissions of SF₆.

The main advantage to the method described here is that it involves a relatively simple conversion of a GC method commonly used for N₂O and SF₆ measurement without compromising N₂O. While laser-based N₂O instruments are emerging, gas chromatography with electron capture detection is commonly used for SF₆ and GC-ECD systems measuring both N₂O and SF₆ will likely be used for the foreseeable future.

2 Methods

The chromatography often associated with the measurement of SF₆ is basically the same as that used to measure N₂O (Elkins 1980; Elkins et al., 1996; Hall et al., 2007). Typically, a 10-port or 12-port, 2-position gas sample valve (GSV) is used to direct an air sample through two columns and an ECD. With calibration and in situ systems we currently use a 12-port GSV (Valco Instrument Co., Houston, TX) operated in “heart cut” mode to prevent the majority of the oxygen/nitrogen in the sample from reaching the ECD (Fig. 1). Upon injection, the sample is directed onto the pre-column. When the gases of interest reach the main column, the GSV is switched, directing flow from the main column to the ECD, while flow through the pre-column is reversed (backflushed) to prevent unwanted compounds from accumulating on the main column. A pressure perturbation results in a slight baseline change, followed by the elution of N₂O and SF₆ (Fig. 2a). We have used this technique with Shimadzu (Kyoto, Japan), Valco,

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small changes in CO₂, and not affected by varying amounts of CO₂ in the sample or by small changes in the pressure of the dopant delivery system. At 2500 ppm CO₂, the sensitivity to +/- 10 ppm fluctuations in CO₂ dopant is a factor of 10 less than the typical 1-sigma precision for both N₂O and SF₆. Insensitivity to sample CO₂ was further
5 verified by comparing air samples with and without CO₂ (CO₂ was completely removed using sodium hydroxide-coated silica (Ascarite)). The response ratios for samples with and without CO₂ was 0.99964 +/- 0.00033 for N₂O and 0.9998 +/- 0.0046 SF₆, which translates into biases of 0.1 ppb and 0.001 pmol mol⁻¹ (ppt), respectively, and is insignificant considering the much smaller natural variations of CO₂ in the atmosphere.

10 The use of CO₂ doping may also minimize interference between N₂O and CO that might be produced from Porapak. Any CO produced would likely react on hot metal surfaces to form CO₂ (Fehsenfeld et al., 1981), which could interfere with N₂O if not for the excess CO₂ supplied by doping. We have not determined if column-generated CO interferes with N₂O when using P-5 carrier gas.

15 One additional change was made involving the column carrier gas head pressure. Electronic pressure controllers (EPC), used to set the head pressures on the main and pre-columns, were installed in late 2005 replacing mass flow controllers. This was done to reduce the effects of changes in room pressure on flow rates. The EPCs are less sensitive to changes in ambient pressure, possibly because they control pressure
20 relative to a reference pressure (ambient pressure in this case) and thus maintain a constant ΔP relative to the ECD outlet.

3 Results

3.1 Calibration instruments

25 With the three-column system, the SF₆ precision was improved from 1–2% to better than 0.5%. SF₆ precision with the three-column system has been consistently better than that of comparable systems using two columns. SF₆ precision determined from

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analysis of various gas standards is shown in Fig. 4. Precision is about a factor of two better than that of a two-column system using a Valco ECD. It is more than a factor of two better than a previous two-column system using an Agilent ECD (ver. 2). The precision of the ver. 2 system was more variable, and seemed to show some improvement in late 2005/early 2006 prior to the modifications. Some of this improvement was due to the installation of the EPCs (late 2005). It is clear that SF₆ precision benefits from stable flow control. However, additional improvements in SF₆ precision can be seen to correspond with the addition of the MS-5A post-column in mid-2006.

A further benefit is that the current system has been essentially maintenance-free over the last four years. SF₆ retention time, peak width, and peak response have remained very stable since the modifications were made. Compared to our original system (ver. 1), the use of N₂ carrier gas has eliminated some of the variability caused by variations in P-5 quality. High-purity nitrogen is more easily obtained, and the quality of N₂ is more consistent than that of P-5, which can sometimes contain significant amounts of SF₆ (Elkins et al., 1996).

The improvements in precision and stability have allowed us to develop an improved reference scale for SF₆ based on standards prepared by gravimetric methods. Nine compressed gas standards were prepared between 2003 and 2005 as described in Hall et al. (2007). These were combined with seven standards prepared in 2000 to establish the NOAA-2006 SF₆ calibration scale (adopted by WMO/GAW) over the range 1-10 ppt (dry air mole fraction) (Table 2) (see also http://www.esrl.noaa.gov/gmd/ccl/sf6_scale.html). The NOAA-2006 scale compares within 0.01 ppt with the Scripps Institution of Oceanography (SIO-05) (Rigby et al., 2010) and within 0.03 ppt of the University of Heidelberg Scale (Hall et al., 2011).

With GC version 3, the standards were analyzed several times relative to a natural-air working standard. The response was linear and a fit of the relative response (peak height) showed a zero intercept. Although a zero intercept was expected from previous work, we were not able to confirm this until the improved precision was realized with GC ver. 3. This important feature means that calibrations can be carried out by ratio

to a single working standard. We are also exploring the range over which linearity can be assumed; currently 1–10 ppt. Reproducibility (1-sigma) of SF₆ assignments has improved from 0.05–0.07 ppt using GC version 1, to 0.03–0.05 ppt using GC version 2, and finally to < = 0.02 ppt using GC version 3.

Five secondary natural air standards are used to verify the instrument response function (calibration run) every other month. The SF₆ response function is stable enough that more frequent calibration is not necessary. This was not true of GC ver. 1. Secondary standards were assigned values on the NOAA-2006 scale. Using the five secondary standards to predict the value of the working standard during each calibration run (linear model (a): $Y = aX$), we find that the predicted value of the working standard has varied by no more than 0.011 ppt over a three-year period. The standard deviation of predicted values is 0.003 ppt. Similar results were achieved when the intercept “b” was not set to zero (linear model (b): $Y = aX + b$). In this case the range of predicted values varied by no more than 0.013 ppt and the standard deviation of all predictions was 0.004 ppt.

Regular analysis of other natural air standards also shows improved precision and stability (Fig. 5). The reproducibility of SF₆ assignments is particularly good following the 2006 modifications. Standard deviations of a “target” standard analyzed between mid-2006 and 2009 are 0.012 and 0.014 ppt using linear models (a) and (b) respectively. Other secondary standards show similar precision on repeated analysis. The long-term precision is comparable to the short-term precision, indicating that there are no significant long-term variations that would lead to a time-dependent bias. Regulator effects might be one reason why SF₆ measurement precision is better for the working standard (blue symbols in Figs. 4 and 5) compared to the target cylinders. We have noticed that some regulators seem to perform better than others, even among the same model number. While long-term conditioning to the air stream (i.e. dedicated regulators) may help in some cases, it does not always lead to improvements.

We have performed limited testing of the current configuration (ver. 3 in Table 1) with an Agilent μ -ECD (G2397A). An Agilent μ -ECD was installed in the calibration GC and

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the column outlet was simply moved from one detector to the other while all operating conditions remained the same. The smaller internal volume of the μ -ECD results in peaks with slightly smaller peak width (see Supplement Fig. S1). While the optimal CO_2 dopant concentration is lower with the Agilent μ -ECD (500–700 ppm) (Andrew Crotwell, personal communication, 2010) compared to the Agilent ECD (\sim 2500 ppm), precision, peak shape, and baseline behavior are similar. While we have not used this technique with a μ -ECD for extended periods of time, limited testing indicates that this technique is fully compatible with the μ -ECD.

3.2 In situ instrument

We have also performed modifications similar to those described above to an in situ N_2O and SF_6 measurement system installed at a mountain site (Niwot Ridge) in Colorado. The original GC configuration was similar to ver. 1 (Valco ECD), but is now operated with a MS-5A post-column and CO_2 -doped N_2 carrier gas similar to ver. 3. Figure 6 shows hourly data from this site for selected periods in 2006 and 2009. SF_6 precision during periods without obvious pollution events was \sim 2% in 2006 and $<$ 0.5% in 2009, a factor of four improvement. Small pollution events can be seen in Fig. 6b, which would have gone undetected earlier.

In addition to the post-column, mass flow controllers used to control carrier gas flows and electrical connections to the ECD were replaced. The improvement in SF_6 precision is due to a combination of these modifications, which were performed simultaneously. We do not know exactly how much the addition of the post-column contributed to the improved precision. Based on results from N_2O , which is measured on the same detector, we estimate that at least one third of the improvement is due to the addition of the post column. The true effect is probably greater than this because N_2O would benefit most from improved carrier gas flow control.

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3.3 A history of atmospheric SF₆

A history of global atmospheric SF₆ was compiled from a combination of grab samples (flasks) and in situ measurements at multiple sites dating to 1995. A history based on two observing programs (one based on flasks and the other in situ) was developed by combining flask and in situ results into a single integrated record at each observing site. Both flask and in situ records offer different advantages and dis-advantages. Flask observations yield greater spatial coverage with lower temporal resolution. Flasks are analyzed on a single laboratory instrument, which should, in principle, be subject to fewer calibration and operational issues than instruments located at remote sites. However, because a single instrument is used, instrument malfunction or performance would affect all flasks analyzed during that period. In situ observations offer greater temporal resolution but lower spatial coverage compared to flasks. Calibration transfer issues would likely be averaged out over time since each in situ instrument is calibrated independently. Each in situ instrument is calibrated using two natural air working standards with concentrations that differ by ~10% to account for non-linearity or non-zero intercepts. These standards are traceable to the NOAA-2006 scale via the primary calibration GC (described above) and are replaced every 1–2 yr.

Figure 7 shows global and hemispheric mean SF₆ from 1995 to present from the integrated SF₆ record. SF₆ data from six in situ sites (BRW, SUM, NWR, MLO, SMO, SPO) were merged with flask data from twelve sites (ALT, SUM, BRW, MHD, NWR, THD, MLO, KUM, SMO, CGO, PSA, SPO) (see Table 3, Supplement Fig. S2). At sites in which both flask and in situ data exist, monthly mean (flask) and median (in situ) were averaged together weighted by standard deviation. Data gaps (typically only a few months) were filled using linear interpolation. Monthly median in situ data were used instead of monthly mean in order to minimize the effects of local influence in the in situ records. Flasks are normally collected under background conditions. In most cases 3–5 flasks pairs per month were combined with ~400–700 hourly in situ samples (data source: <http://www.esrl.noaa.gov/gmd/hats/combined/SF6.html>).

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The history was extended to the mid-1980's by analyzing cylinders of air collected at Niwot Ridge, Colorado. The archive consists mostly of 29.5-L aluminum cylinders collected specifically as archive samples to be stored, or used as working standards and not completely exhausted. Archive samples were screened for background conditions by examining trace gases such as CFC-12 as well as local wind direction at the time of sampling. Eight out of 68 potential archive samples were rejected based on concentrations of CFC-12, CFC-11, and/or wind direction not having been consistent with background air masses.

The integrated data set consists of both high-frequency in situ measurements and low frequency flask measurements. The uncertainty (standard error) for each month at each site was calculated using an effective number of independent measurements (N_{eff}). Although there are as many as 744 individual measurements per month at some sites, not all measurements are independent. We estimated the number of independent samples by calculating the time scale of auto correlation of hourly samples. Typically, hourly samples are auto-correlated ($\rho > 0.1$) for 12–48 h at a time at each site. This seems to correspond to the timescales of diurnal cycles or synoptic wave activity. Thus, we estimate that a monthly median estimate of SF₆ concentration consists of ~15 independent estimates from in situ data and 3–4 independent estimates from flasks. Monthly standard deviations for each site were scaled by the square root of N_{eff} . The monthly uncertainty at each site was then determined as the mass-weighted mean standard error from each program (flasks and in situ) when data from both programs were available. An additional uncertainty was introduced when the mean mole fraction from each program differed by more than the combined standard errors. Hence, differences between two independent systems are incorporated as an additional term in the uncertainty. The mean difference between global mean SF₆ computed using flask and in situ data from 2000–2010 is < 0.01 ppt (see Fig. S3).

Monthly data were averaged in a mass-weighted (cosine latitude) fashion across seven latitude bins to produce global and hemispheric means. Uncertainty in the global mean was estimated by computing the global mean numerous times (Monte-Carlo

method), randomly adjusting the monthly mean at each station within a $2\text{-}\sigma$ window (Normal distribution). Further, data gaps (up to 30 % of the data) were introduced at random to test the robustness of the gap-filling interpolation method. The integrated global mean data record shown in Figure 7 was derived from 500 simulations.

5 Global mean SF_6 from 1985–2005 can be approximated by the function $y = 0.125 \pm (0.012) + 0.215 \pm (0.001) \times (t - 1985)$. This function supersedes that of Geller et al. (1997), and results from an updated SF_6 calibration scale, reanalysis of the archive samples analyzed by Geller et al. (1997), improved measurement precision, and a longer data record.

10 3.4 SF_6 growth rates

Global monthly mean SF_6 concentrations were used to derive global SF_6 growth rates by subtracting global monthly mean concentrations in successive years (e.g. January 2003–January 2002, February 2003–February 2002, etc.) (Fig. 8). Because SF_6 is extremely long-lived in the atmosphere, the increase in SF_6 concentration from one year to the next is essentially proportional to the emissions in that year. A close approximation of emissions in Gg yr^{-1} can be obtained by multiplying the growth rate (ppt yr^{-1}) by 25 (Maiss and Brenninkmeijer, 1998).

15 The average growth rate from 1986–2005, determined from N.H. archive samples was $0.214 \text{ ppt } \pm 0.001 \text{ yr}^{-1}$. In agreement with Levin et al (2010) and Rigby et al (2010), our results show that the global growth rate of SF_6 (emissions) increased gradually between 2001 and 2006, and rapidly from 2006–2008, reaching a peak of $0.30 \pm 0.03 \text{ ppt yr}^{-1}$ (or $\sim 7.4 \text{ Gg yr}^{-1} \text{ SF}_6$) in 2007–2008. SF_6 emissions increased at an average rate of $5\% \text{ yr}^{-1}$ from 2001–2008. This corresponds to a period of rapid growth in world gross domestic product (GDP). The general features of global SF_6 growth rates since 1995 derived from these data are consistent with those of Levin et al. (2010) and Rigby et al. (2010).

25 Some inter-annual variability in SF_6 emissions is suggested from this simple analysis. Some of this apparent inter-annual variability is probably due to atmospheric

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(Fig. 10b) except during 2008–2009, during which no large transport-driven declines are observed.

The observed decline in the SF₆ growth rate from the peak in 2008 through mid-2009 is 14 %. From modeled SF₆ (constant emissions) and other long-lived tracers, we estimate that atmospheric transport is responsible for at most ~30 % of the observed change. Therefore, the change in SF₆ emissions during 2008–2009 is estimated to be ~10 %. This decline also coincides with a slowing of the world economy, with GDP decreasing ~0.3 % following a decade of growth. Although such a rapid decline in emissions might seem unlikely, it is not unprecedented. Global SF₆ emissions declined 27 % from 1995 to 1998 (Maiss and Brenninkmeijer, 2000) and 8 % from 1996–1997 (Levin et al, 2010), and increased 8 % between 2006 and 2007. The decline in SF₆ emissions in 2008–2009 appears to be short-lived, as SF₆ growth rates began to increase in early 2010, approaching their peak value observed in 2008.

4 Conclusions

By modifying a common GC technique used to measure N₂O and SF₆, we have improved the precision of SF₆ measurement by a factor of 2–3. The precision associated with a calibration instrument, used to transfer the NOAA-2006 scale to tertiary compressed gas standards, has been improved in recent years and is currently around 0.3 % or ~0.02 ppt based on ambient SF₆ levels. As this technique is extended to instruments currently in use, our ability to quantify SF₆ emissions on global and regional scales should improve.

Mixing ratios of SF₆, measured in flasks collected at 12 sites around the world, and measured in situ at six sites, continue to increase in the troposphere. The growth rate (and inferred emissions of SF₆) has increased in recent years reaching a maximum of 0.30 ± 0.03 ppt yr⁻¹. This is almost 40 % larger than the average growth rate of 0.21 ppt yr⁻¹ deduced from archive samples in the N.H. collected between 1987 and 2005. When SF₆ growth rates are examined on a station-by-station basis, all N.H. sites

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show two similar features: an increase in the growth rate from 2005–2008, and a decrease in the growth rate from 2008 through mid-2009. The increase in SF₆ growth rate corresponds to an increase in World GDP through 2008, while the recent slowing of SF₆ growth rate corresponds to a decline in world economic growth. The short-lived decline in SF₆ emissions observed during 2008–2009 is estimated to be about 10 % of the 2008 maximum.

Supplementary material related to this article is available online at:
<http://www.atmos-meas-tech-discuss.net/4/4131/2011/amtd-4-4131-2011-supplement.pdf>

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Table 1. Configuration of three SF₆ GCs used for calibration since 1999.

	Version 1	Version 2	Version 3
Period of use	Sept. 1999–Feb. 2003	Feb. 2003–Jun. 2006	Jul. 2006–present
Pre-column	2 m × 3.7 mm I.D., 4.76 mm O.D. Porapak-Q, 80/100 mesh	2 m × 3.7 mm I.D., 4.76 mm O.D. Porapak-Q, 80/100 mesh	1 m × 3.7 mm I.D., 4.76 mm O.D. Porapak-Q, 80/100 mesh
Pre-column <i>T</i> (°C)	55	56	90
Main column	3 m × 3.7 mm I.D., 4.76 mm O.D. Porapak-Q, 80/100 mesh	3 m × 3.7 mm I.D., 4.76 mm O.D. Porapak-Q, 80/100 mesh	2 m × 3.7 mm I.D., 4.76 mm O.D. Porapak-Q, 80/100 mesh
Main-column <i>T</i> (°C)	55	56	90
Post-column	NA	NA	0.91 m × 3.18 mm O.D. molecular sieve 5A, 40/60 mesh
Post-column <i>T</i> (°C)	NA	NA	185
Flow control	MFC	MFC/EPC	EPC
Detector	Valco (140BN)	Agilent (G1533A)	Agilent (G1533A)
Carrier gas	5 % CH ₄ /Ar (P-5)	N ₂ (w/CO ₂ doping)	N ₂ (w/CO ₂ doping)
SF ₆ retention time (s)	~ 440	515–545	305
SF ₆ peak width (s)	~24	30	23
N ₂ O retention time (s)	~ 345	405–435	410
N ₂ O peak width (s)	~18.5	23	27
ECD <i>T</i> (°C)	350	340	340/370
Run time (s)	1800	900	720
GSV switch (s)	240	300–320	180
CO ₂ retention time (s)	~270	~365	~475

T: temperature; MFC: mass flow controller; EPC: electronic pressure controller.

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Table 2. Standards that define the NOAA-2006 SF₆ calibration scale.

Cylinder	Year	Prepared (ppt)	Assigned (ppt)	Residual (ppt)
FA-1861	2000	2.41	2.39	-0.02
FA-1878	2000	2.91	2.89	-0.02
FA-1843	2000	4.73	4.76	0.03
FA-1850	2000	4.51	4.53	0.02
FA-1851	2000	4.32	4.35	0.02
FA-1856	2000	1.14	1.15	0.01
FA-1865	2000	4.12	4.13	0.01
FA-2205	2000	4.94	4.92	-0.02
FA-2207	2003	5.98	5.98	0.00
FA-2208	2003	6.97	7.01	0.03
FA-1940	2005	1.52	1.49	-0.03
FA-2139	2005	3.13	3.14	0.01
FA-2557	2005	3.86	3.85	-0.01
FA-2567	2005	5.99	6.02	0.03
FA-2569	2005	7.92	7.90	-0.02
FA-2585	2005	9.83	9.79	-0.04

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Table 3. Sampling sites.

Sampling site	Abbr.	Lat., Long.	Alt. (masl)	Obs.
Alert, Nunavut	ALT	82.5, -62.5	54	flask
Summit, Greenland	SUM	72.6, -38.5	3238	in situ
Barrow, AK	BRW	71.3, -156.6	30, 30	flask, in situ
Mace Head, Ireland	MHD	53.3, -9.9	25	flask
Trinidad Head, CA	THD	41.1, -124.2	107	flask
Niwot Ridge, CO	NWR	40.0, -105.5	3523, 3025	flask, in situ
Cape Kumukahi, HI	KUM	19.5, -154.8	3	flask
Mauna Loa, HI	MLO	19.5, -155.6	3397, 3397	flask, in situ
Cape Matatula, Am. Samoa	SMO	-14.2, -170.6	77, 77	flask, in situ
Cape Grim, Australia	CGO	-40.7, 144.7	94	flask
Palmer Station, Antarctica	PSA	-64.9, -64.0	10	flask
South Pole, Antarctica	SPO	-90.0, -24.8	2810, 2810	flask, in situ

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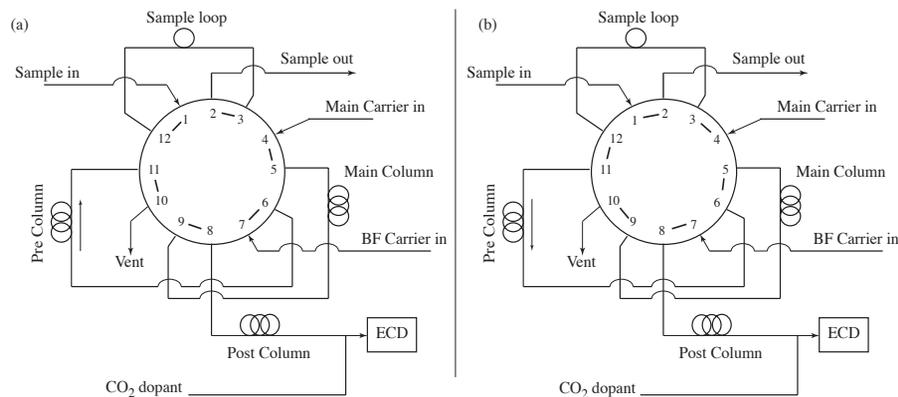


Fig. 1. Diagram of 12-port gas sample valve (GSV) in (a) load, and (b) inject modes. Note that Air (O₂/N₂) is vented during inject mode, and the sample is directed from the main and post columns to the ECD in load mode.

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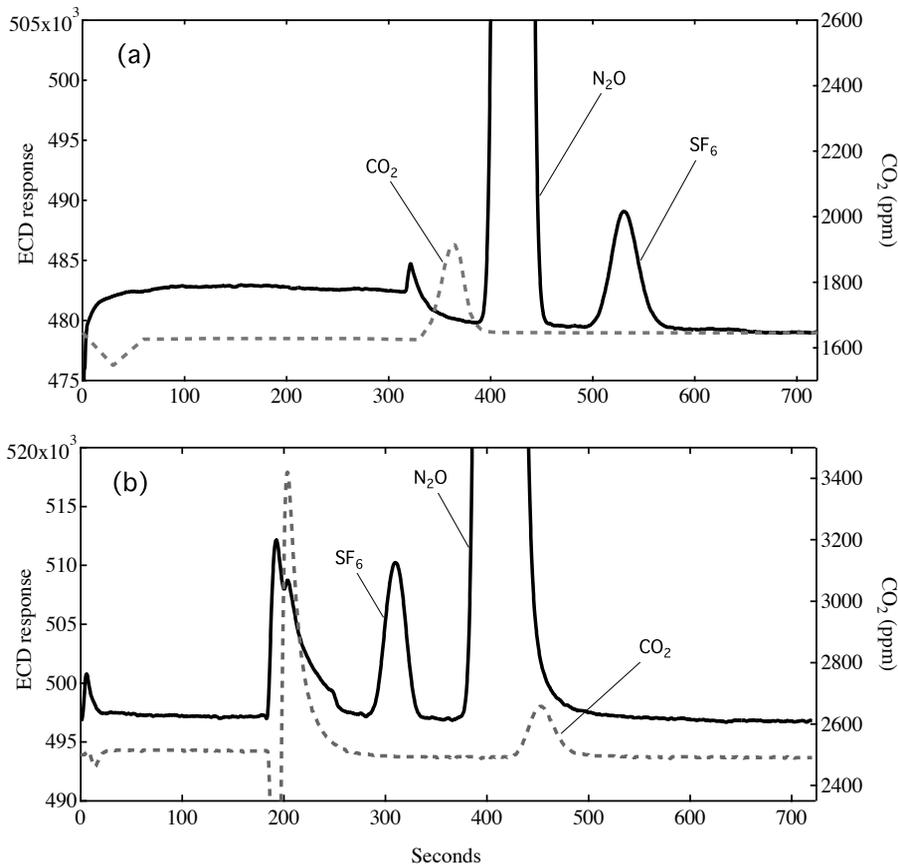


Fig. 2. Sample chromatograms from two-column N₂O/SF₆ analysis system (upper panel, version 2) and three-column system (lower panel, version 3) (see Table 1). CO₂ was measured at the outlet of the ECD.

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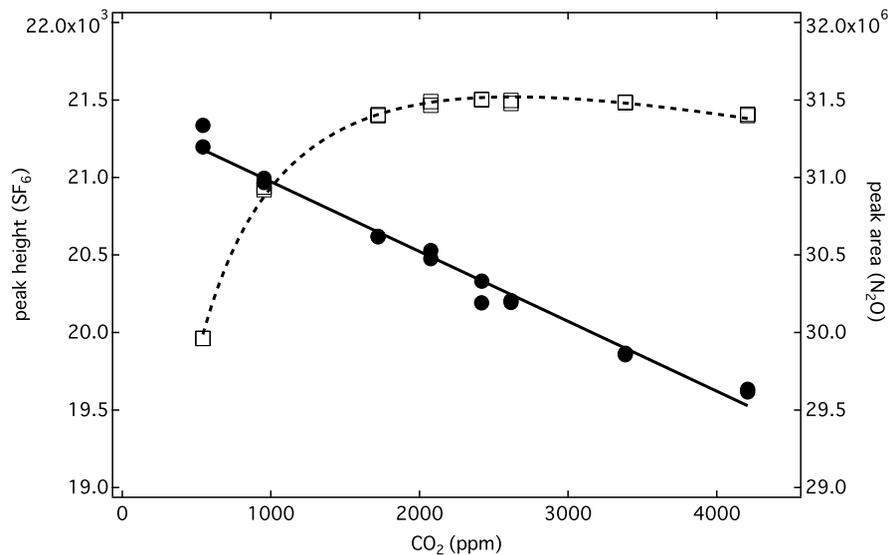


Fig. 3. Effect of CO₂ on SF₆ (left axis, closed circles) and N₂O (right axis, open squares) response (arbitrary units).

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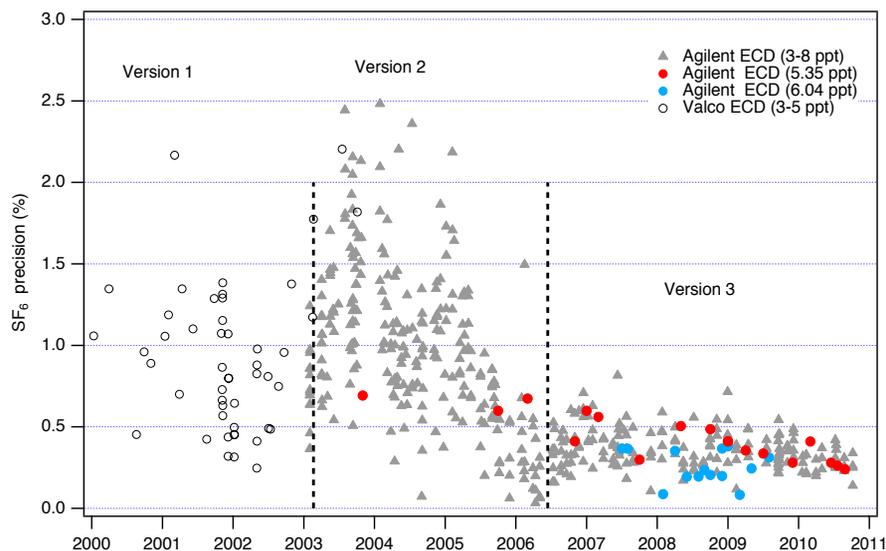


Fig. 4. History of SF₆ precision (%) from GC-ECD systems used for calibrations.

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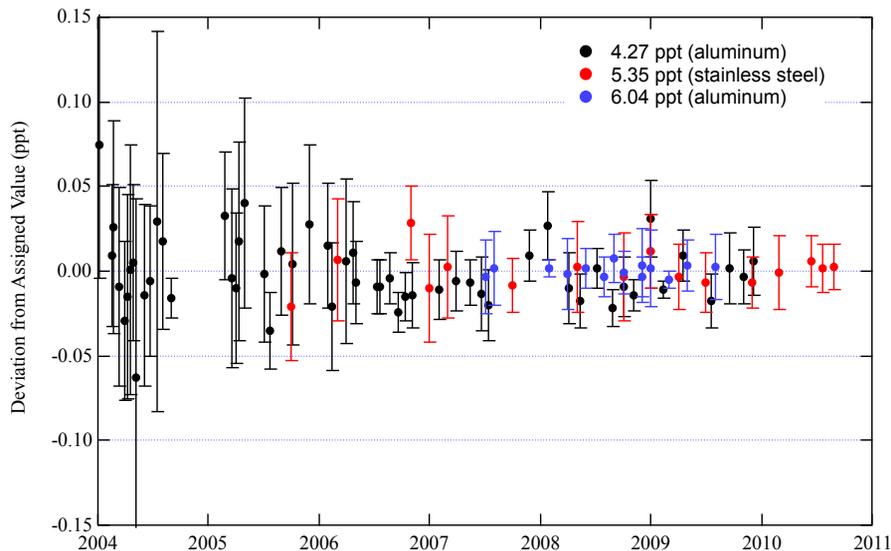


Fig. 5. History of variability in SF₆ calibration assignments. Since 2006 the mean deviation has been -0.005 ppt, 0.002 , and 0.005 ppt for 4.27, 5.35, and 6.04 ppt target standards, respectively.

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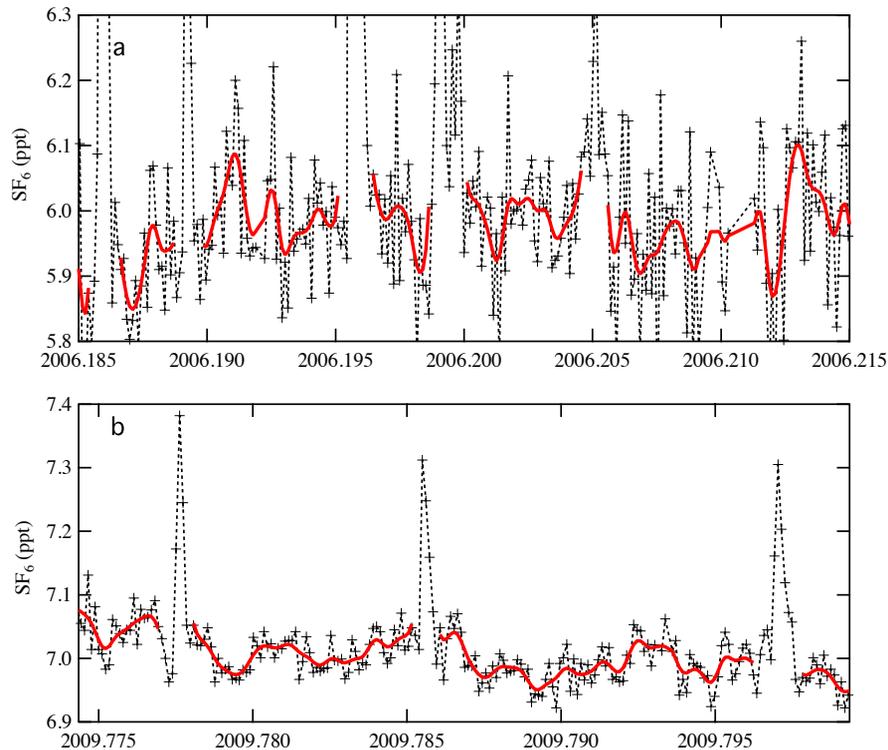


Fig. 6. Time series of SF₆ at Niwot Ridge, Colorado showing baseline conditions and pollution events during two periods: **(a)** prior to GC modification; **(b)** following GC modification. Both panels show a smooth fit to baseline data (approximate) (red lines).

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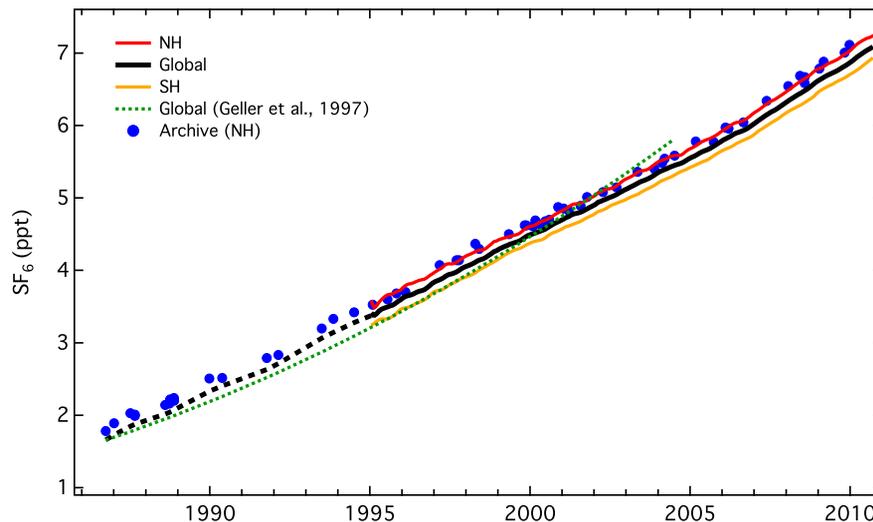


Fig. 7. Global and hemispheric mean SF₆ from flask and in situ data along with NH archive samples. The dashed line prior to 1995 represents the global mean derived from NH archive data (computed from a lowest fit to the NH archive record minus the mean difference (0.14 ppt) between NWR and the global mean from 1995–2005). A linear fit of global mean SF₆ as a function of time (t) from 1986–2005 yields: $y = 1.25 (0.01) + 0.215 (0.001) \times (t - 1985)$.

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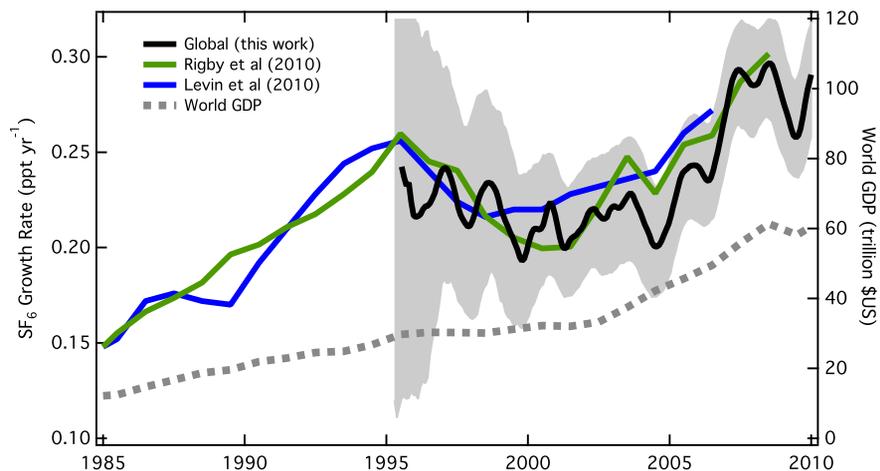


Fig. 8. Global growth rate (ppt yr^{-1}) determined from flask and in situ combined SF₆ dataset. (shaded region is 2σ). Also shown are growth rates derived from emission estimated of Levin et al. (2010) and Rigby et al. (2010) along with World GDP (World Bank, 2011).

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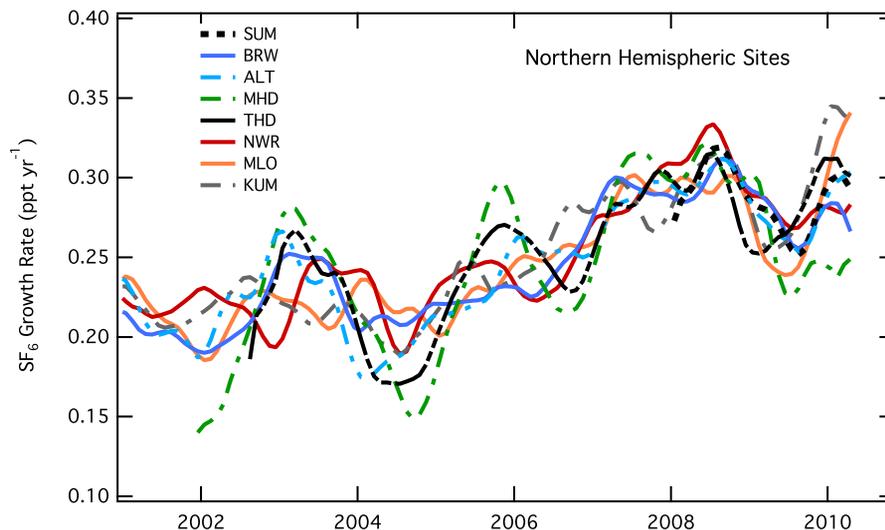


Fig. 9. SF₆ growth rates calculated on a site-by-site basis at eight N.H. sites.

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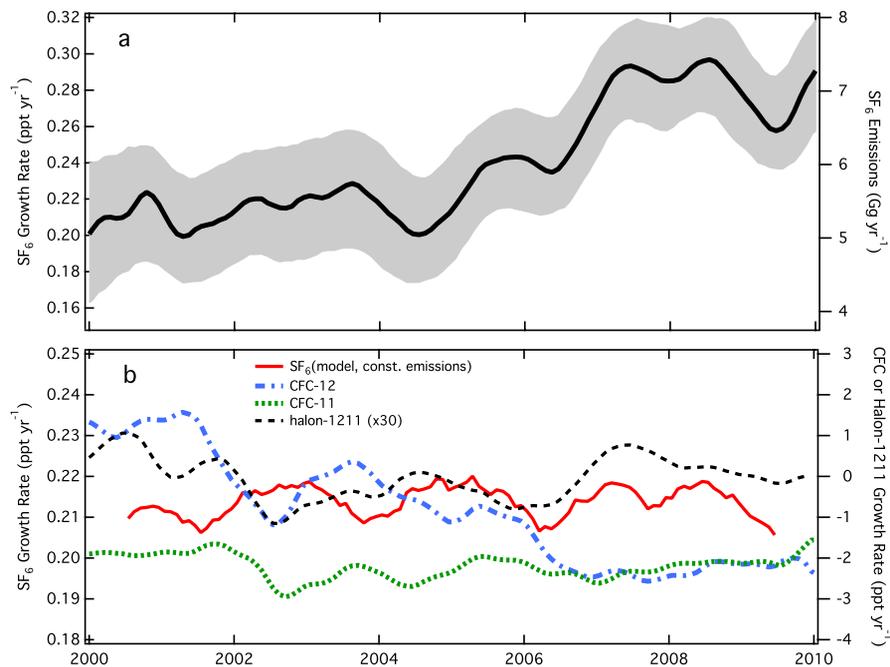


Fig. 10. (a) SF₆ growth rate derived from global mean SF₆ (shaded region is 2σ), along with corresponding emissions (Gg yr⁻¹) calculated with a 2-box model (Geller et al., 1997); (b) observed growth rates of CFC-12 (blue), CFC-11 (green), and halon-1211 (black, detrended) derived from global mean data, along with SF₆ growth rate derived from model results at four sites using constant SF₆ emissions (solid red) to assess transport-driven changes in the SF₆ growth rate.

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