

Dear Editor:

Thank you for allowing us to revise and improve our manuscript. We are thankful to reviewers for their critical comments and constructive feedbacks on our manuscript. We have done our best to incorporate all the comments into our manuscript in this revised version. We believe the reviewer's input greatly improved the quality of our manuscript. Below is our detailed responses. Reviewers' comments are in blue and our responses are in black. The differences between the submitted and revised manuscript are marked for reading convenience, and the pdf files including supplement are attached at the end of this file.

Jin Ma (On behalf of co-authors)

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Response to Review #1

J. Elliott Campbell (Referee)

We would like to thank Professor Campbell for his time and effort to read our paper and evaluate our research.

Major Comments

To what extent can you discuss and analyze the importance of implementing CS₂ as a separate tracer? Previous studies assume CS₂ emissions convert on emission and only have a single OCS tracer. I wonder if one region this may be important is the Asian anthropogenic outflow. In previous studies without the separate tracers, the high mixing ratios should be more immediately over Asia while in your TM5 simulations the highest mixing ratios should be somewhat downstream of the Asian source. Perhaps a feature like this can be seen in TES data. It may be relevant to draw comparisons to related studies with CO₂ such as: Suntharalingam, P., Randerson, J. T., Krakauer, N., Logan, J. A. and Jacob, D. J.: Influence of reduced carbon emissions and oxidation on the distribution of atmospheric CO₂: Implications for inversion analyses, *Global Biogeochem. Cycles*, 19, GB4003, 2005.

The inclusion of CS₂ in our model gives the possibility to validate the geophysical distribution and vertical profiles in the future. It is also worth to note that CS₂ has a different source distribution compared to COS. E.g., CS₂ direct emissions are mainly from the industrial production of rayon. Another issue is the atmospheric lifetime of CS₂. As referee #2 also discussed, the lifetime of CS₂ is more likely shorter than what we used in the current implementation (Kahn et al., 2017). It is more likely less than 4 days, and we will test the sensitivity of the CS₂ lifetime in TM5. In the revised manuscript, we will include CS₂ sensitivity tests by varying the lifetime from ~6 days (current setting), to 3 days (Kahn et al., 2017). Another sensitivity test that we will report on is the instantaneous transfer of CS₂ to COS, and compare it with the current model settings.

In the abstract, the authors find that the missing source shows little inter-annual variation but large seasonal variation. While figure 6e provides some information on this seasonality, further plots and discussion would be helpful to explore this seasonality. Consider adding maps to the supplement of the optimized fluxes from 6e for 4 seasons and time series of regional averages.

Thanks for the suggestion. In order to better investigate the inter-annual variations of the missing source, we have done further time-series analysis for regions. Globally, the COS unknown posterior flux is shown, together with the trend, the seasonal signal and the residual (Figure 1). It can be seen that, globally, the seasonal signal is highest during NH spring and lowest during NH fall. The global flux was then split into 8 regions (Figure 2), and the regional COS unknown flux analyzed for these regions is shown in Figure 3. The region NH1 (North America plus part of Pacific and Atlantic Oceans, orange) shows a negative “unknown” flux, indicating that more sinks are needed. This likely points to an underestimation of the biosphere uptake in the prior, since this region (that is well constrained by observations) depicts a clear seasonal cycle in the optimized “unknown” flux. NH2 (Europe, green) and NH3 (Asia, red) have almost the same trend and seasonality, perhaps because they are difficult to separate using the available observations. Tropical regions TR0-3 have similar trend and seasonality, and generally shows a positive flux signal, with little seasonal cycle. This could represent and oceanic signal (underestimated emissions of COS or COS precursors in the prior), a signal

from biomass burning, or an overestimated biosphere sink. The ocean-dominated region SH (blue) has a near neutral flux, with a seasonal cycle that shows higher emissions in local fall and early winter.

We will add some of these analyses in the supplement or in the main text of the revised manuscript.

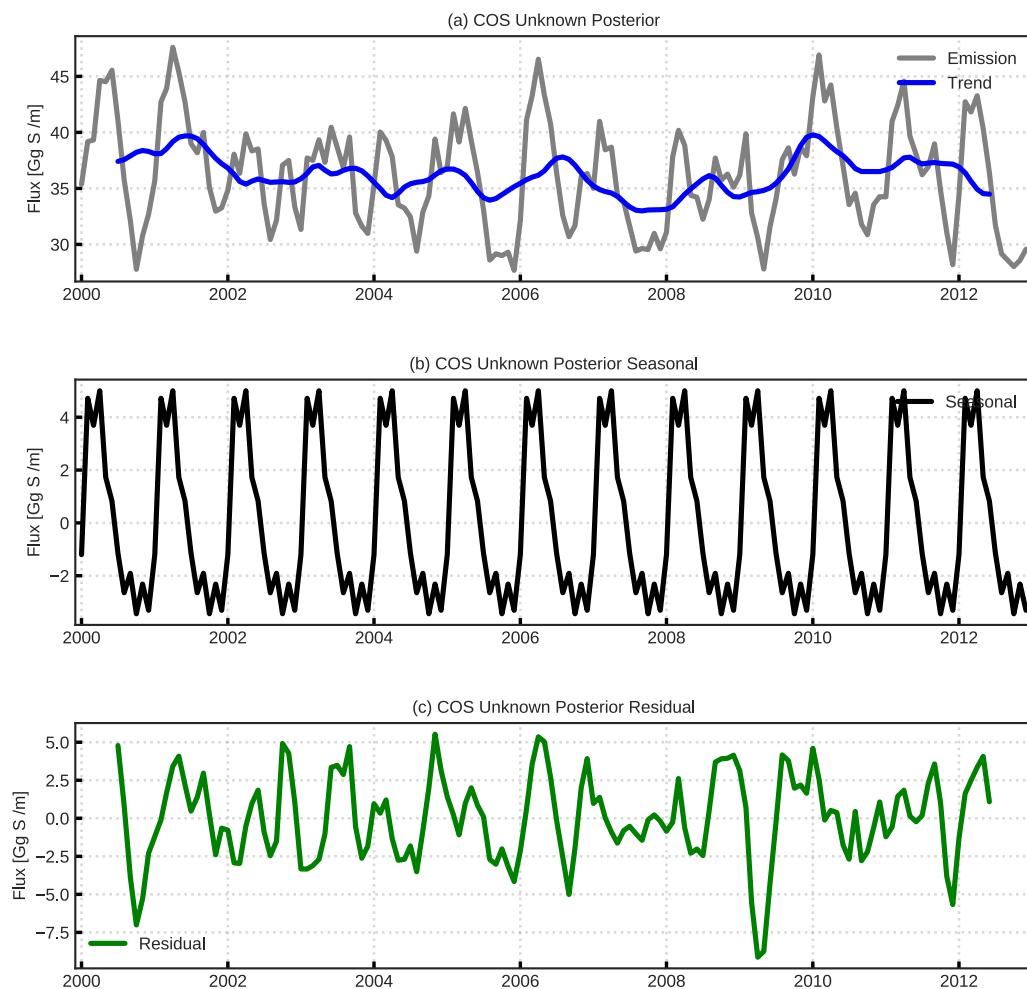


Figure 1. COS posterior flux of the optimized category "unknown" in inversion Su, with its seasonal signal and residual for the years 2000–2012.

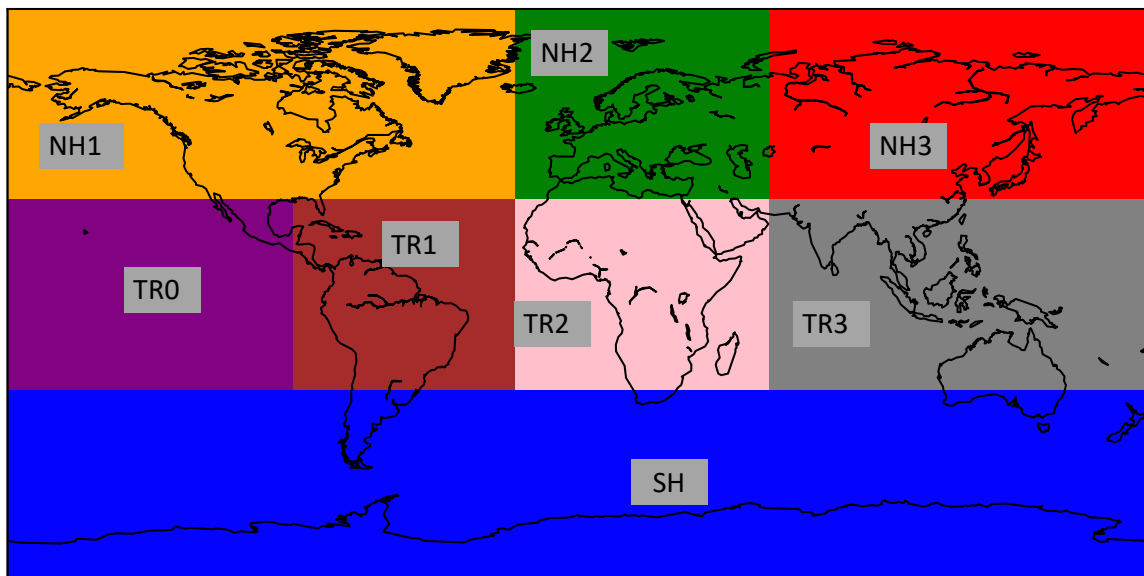


Figure 2. The regional map for analysis of COS fluxes. NH1-3 are areas in Northern Hemisphere. TR0-3 are regions in Tropics. SH refers to the Southern Hemisphere.

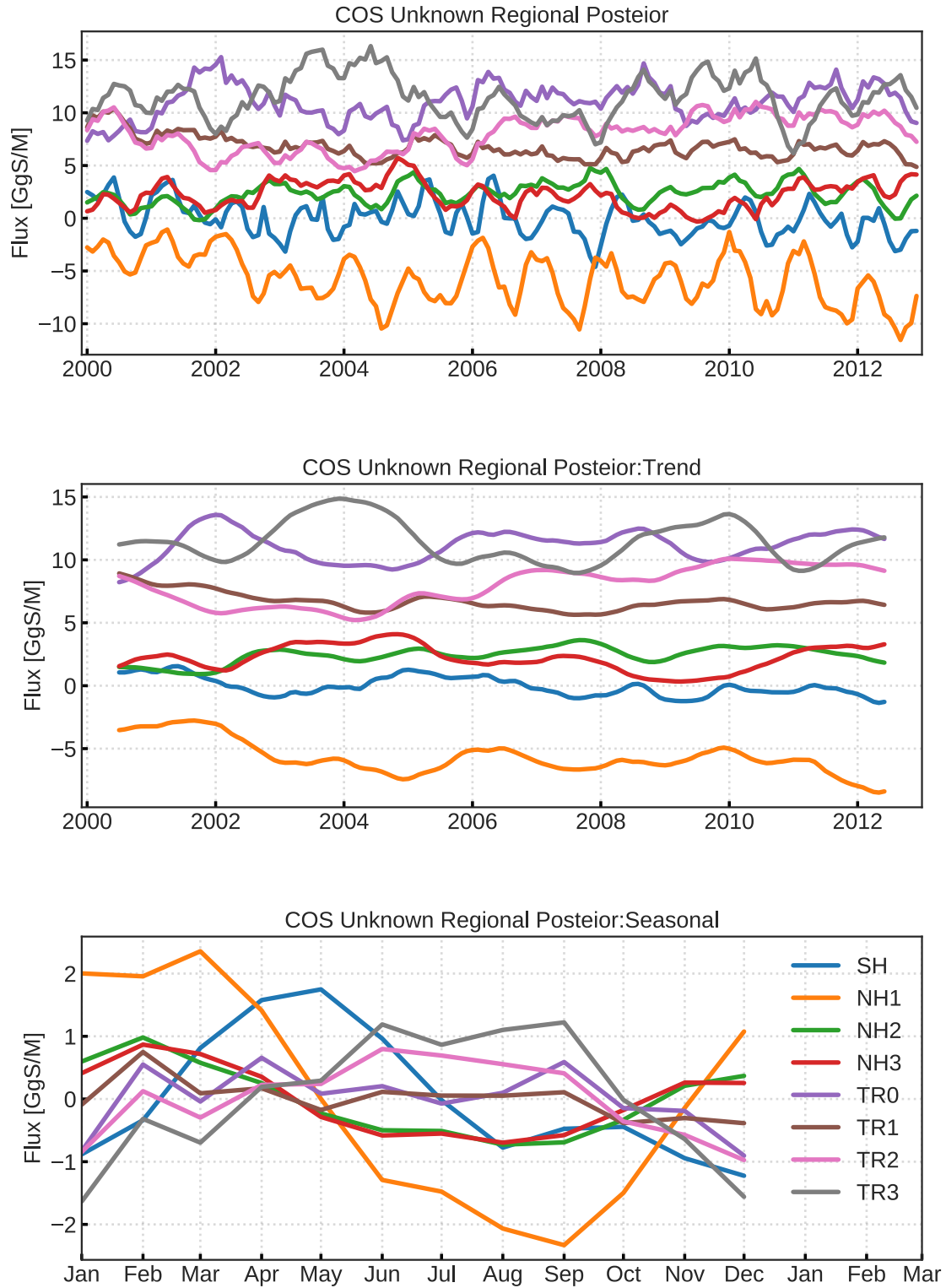


Figure 3. The regional signals of the optimized COS posterior flux (inversion S_u). The upper panel shows flux contribution in each region, and the middle panel shows the trend by filtering out the seasonal and residual signals. The lower panel is the mean seasonal cycle of the COS posterior flux in each region. The colors of the lines are the same as in Figure 2.

The abstract notes that an overestimated sink cannot be ruled out but the manuscript notes that tropical land constraints are not available. This note about missing tropical continental boundary layer data should be added to the abstract for clarity.

We will add this message to the abstract.

Can the authors reconsider this statement about the overestimate sink by further use of the MIPAS data? I don't think you need another inversion run but just a few plots to compare the S1 and S3 runs to the geographic variability in MIPAS. Note that the MIPAS data clearly show global minimums in the convective outflow of the Amazon? I don't think Fig10 is sufficient to explore the MIPAS constraint because the critical dimension is longitude. In the MIPAS tropics there is high mixing ratios in the western tropical Pacific and low mixing ratios over Amazon/Congo. Maps of TM5 at high altitude along with maps of MIPAS are needed to test the validity of the large changes in the biosphere flux from inversions S1 and S3.

This is a good suggestion. We indeed have the feeling that inversion S3 projects the total missing source on the biosphere, and this should impact the longitudinal MIPAS comparison. We will consider some extra figures in the Supplement.

Section 2.2.3 could use some additional explanation with respect to: "The SiB4 model was constrained by a prescribed COS mole fraction of 500 pmol mol⁻¹ outside of canopy." This 500 pmol mol⁻¹ is a placeholder. The actual boundary layer mixing ratio is lower and this SiB flux is best implemented in a model using a first-order dependency on ambient levels. Thus the 1053 GgS/y is likely an overestimate which is consistent with the lower flux reported in Berry et al of 738 GgS/y in which the first-order relationship is used. I think its fine that this study uses a zero-order approach but I think this should be carefully distinguished in the methods from future work that will need to implement the first-order approach. Furthermore it could be noted later in the abstract when the correction is made to obtain 851 GgS/y that this result is closer to the Berry et al result of 738 GgS/y.

We realize that the SiB4 model uses 500 ppt merely as a placeholder, leading to a large uptake. We agree that our current manuscript presents this too much as a deficiency of the SiB4 model, and will adjust this in the discussion, section 2.2.3, and the abstract. The preferred (but more difficult) implementation of a first-order removal in the 4DVAR framework will be addressed in a future study.

Minor Comments

Maybe adjust the abstract wording to slightly improve clarity that plant sink in the TM5 runs are zero order (not first order). For example, "We finally find that the biosphere flux dependency on surface COS mole fraction (which was not modeled in this study) may substantially. . ."

This will be fixed in the manuscript.

The introduction notes Suntharalingam et al. (2008) study which attempted to fit background data by increasing the plant sink but you may also want to reference the Campbell et al. (2008) finding that this upward revision could be validated using direct observations from the continental boundary layer from the intensive INTEx-NA airborne campaign. Campbell, J.

Elliott, et al. "Photosynthetic control of atmospheric carbonyl sulfide during the growing season." *Science* 322.5904 (2008): 1085-1088.

We thank the reviewer to point out the publication. This will be included in the manuscript.

The authors site Lennartz et al. (2017) for the bottom-up ocean emissions but please also note the upward revision in Lennartz et al. (2019). Lennartz, S. T., von Hobe, M., Booge, D., Bittig, H. C., Fischer, T., Gonçalves-Araujo, R., Ksionzek, K. B., Koch, B. P., Bracher, A., Röttgers, R., Quack, B., and Marandino, C. A.: The influence of dissolved organic matter on the marine production of carbonyl sulfide (OCS) and carbon disulfide (CS₂) in the Peruvian upwelling, *Ocean Sci.*, 15, 1071–1090, <https://doi.org/10.5194/os-15-1071-2019>, 2019.

We thank the reviewer to point out the publication, which will be included in the revised manuscript.

Section 2.1.1 should note the lack of observatories in the tropical continental boundary layer. Regarding section 2.2.1, how did you divide the anthropogenic emissions into direct COS and indirect CS₂? Did the Zumkehr data file present emissions separately for direct and indirect? If they didn't then how did you back this out? You may want to look at the emission inventory in Campbell et al (2015) which does present separate emission estimates for direct COS and indirect CS₂. The Zumkehr approach was an extension of Campbell et al 2015.

Campbell, JE, Whelan, ME, Seibt, U, Smith, SJ, Berry, JA, and Hilton, TW (2015), Atmospheric carbonyl sulfide sources from anthropogenic activity: Implications for carbon cycle constraints. *Geophys. Res. Lett.*, 42, 3004– 3010. doi: 10.1002/2015GL063445.

We implemented anthropogenic emissions largely based on Zumkehr et al. (2018) and Campbell et al 2015. We separated the COS emission from the Zumkehr et al. (2018) work according to Table 1 in Lee and Brimblecombe (2016). In Table 1 the authors reported a detailed emission budget for COS and CS₂ in anthropogenic categories. Then we used the ratio of this budget to roughly estimate the direct and indirect COS anthropogenic emissions. In this way we were able to separate the COS and CS₂ direct anthropogenic emissions, which should be correct within the uncertainties. We will outline the followed procedure more clearly in the revised manuscript.

Section 2.2.1 discussed uncertainties in molar yield. It might be worth noting that the uncertainty in the anthropogenic inventory is much larger than the uncertainty in molar yield.

We will clarify this better in the revised manuscript. With more measurements available, we could try to reduce these uncertainties using our inverse modelling framework.

Section 2.2.2 might want to draw comparison of this studies results to previous estimates from biomass burning in Campbell et al (2015) and open burning in Stinecipher et al. (2019).

We will mention a few differences in the revised manuscript. The major difference is that we considered new biofuel emission factors in South Asia.

Regarding the poor posterior fit at NWR and THD, are there references in the CO₂ inversion literature that had the same difficulty? These sites are designed to capture background mixing ratio but sometimes they suffer from local influence which would be one reason for the poor posterior fit. One helpful paper you may want to reference is Riley et al.

Riley, W. J., Randerson, J. T., Foster, P. N., and Lueker, T. J. (2005), Influence of terrestrial ecosystems and topography on coastal CO₂ measurements: A case study at Trinidad Head, California, *J. Geophys. Res.*, 110, G01005, doi:10.1029/2004JG000007.

We thank the reviewer to point out the CO₂ paper concerning THD. It is true that THD as discussed in the paper of Riley et al. (2005) is more affected by the local coastal effect and the diurnal cycle of CO₂ fluxes. The current TM5 inverse modelling effort applies a coarse resolution of $6^{\circ} \times 4^{\circ}$ globally, and thus the effect of coastal meteorology at THD is not well captured. Another point is that biosphere flux of COS is applied on a monthly-average basis, and does not account for a diurnal cycle. The NWR station is probably also affected by local land effects that are not well resolved in the coarse simulations.

To compare with CO₂ inversions, we have investigated CarbonTracker North America data and found similarities between COS and CO₂ inversions. For example, at THD, CO₂ has relatively large model-data mismatches of ~ 11 ppm in Summer (Figure 4). For NWR, the CO₂ model-data mismatch is about 3.37 ppm in Summer (Figure 5). In comparison, at MHD the CO₂ model-data mismatch is only 2.04 ppm in Summer inferred from Flask observations. Note that CarbonTracker North America employs a resolution of $1^{\circ} \times 1^{\circ}$ degree, compared to $6^{\circ} \times 4^{\circ}$ degree in our COS study.

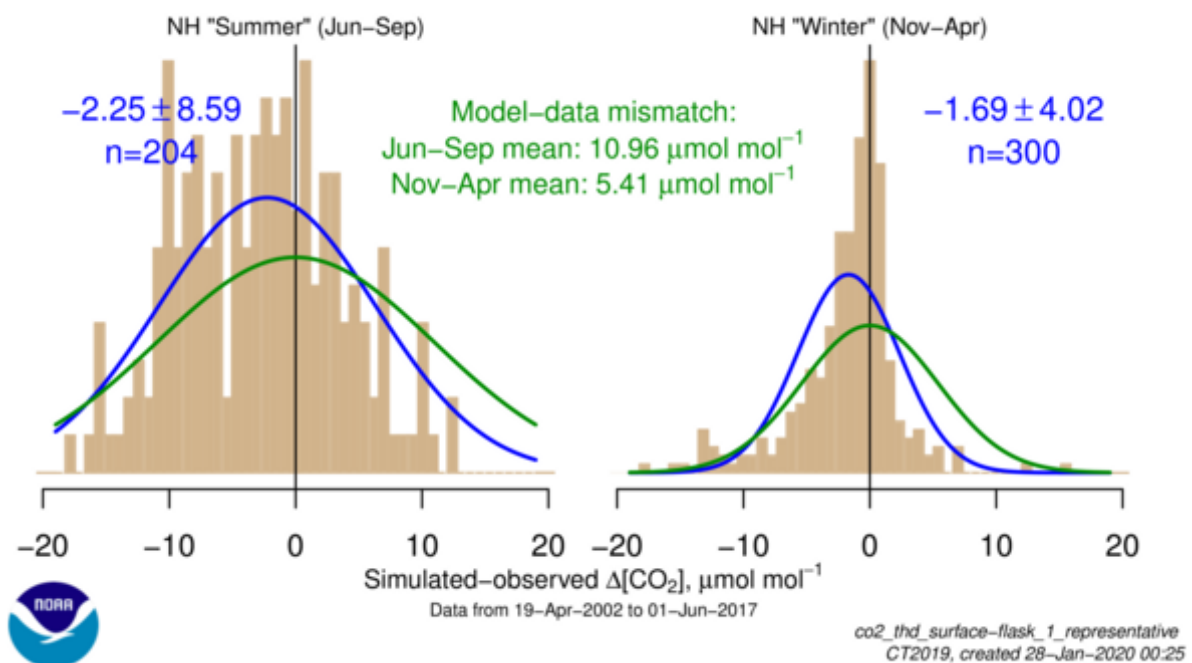


Figure 4. CO₂ observation and simulation by CarbonTracker at THD (source: https://www.esrl.noaa.gov/gmd/ccgg/carbontracker/co2tser.php?ds=co2_thd_surface-flask_1_representative&ed=assim&lastds=co2_nwr_surface-flask_1_representative).

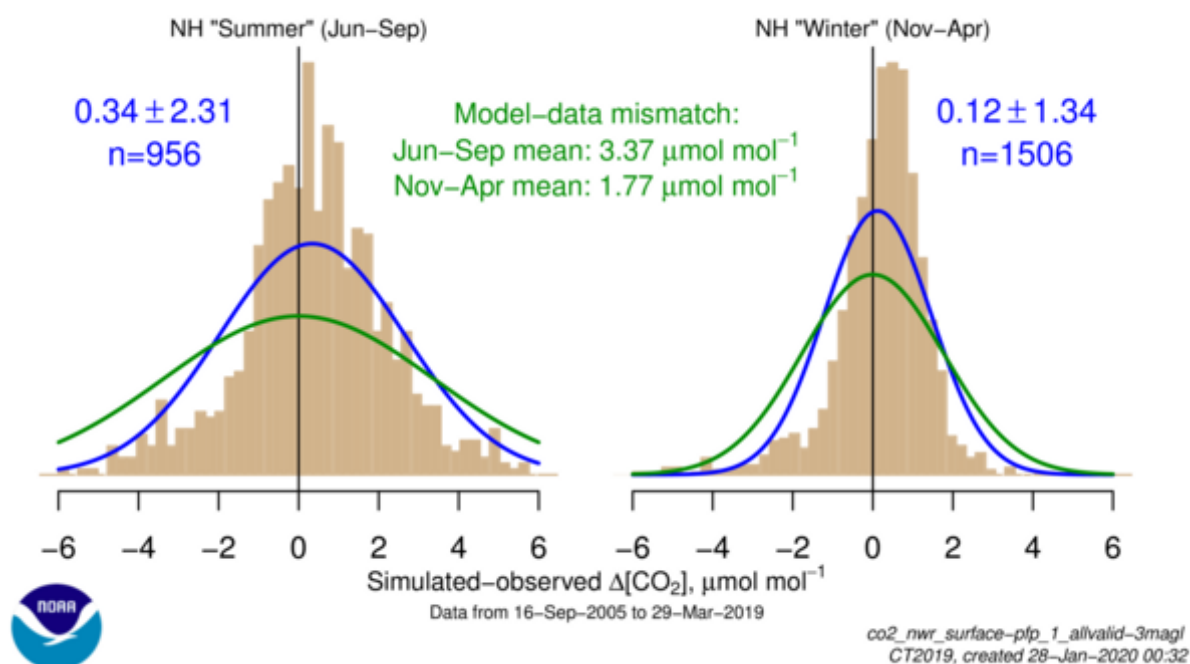


Figure 5. CO₂ observation and simulation by CarbonTracker at NWR (source: https://www.esrl.noaa.gov/gmd/ccgg/carbontracker/co2tser.php?ds=co2_nwr_surface-plp_1_allvalid-3magl&ed=assim&lastds=co2_nwr_surface-insitu_3_nonlocal).

Line 421: “However, observations clearly show a large drawdown of COS near the surface (Hilton et al., 2017; Spielmann et al., 2020).” You may want to reference the INTEX-NA data which is an intensive sampling of vertical profiles in the continental boundary layer (Campbell et al., 2008)

We will include the reference.

References:

1. Khan, A., Razis, B., Gillespie, S., Percival, C., & Shallcross, D. (2017). Global analysis of carbon disulfide (CS₂) using the 3-D chemistry transport model STOCHEM. *Aims Environ. Sci.*, 4, 484-501.
2. Zumkehr, A., Hilton, T. W., Whelan, M., Smith, S., Kuai, L., Worden, J., & Campbell, J. E. (2018). Global gridded anthropogenic emissions inventory of carbonyl sulfide. *Atmospheric Environment*, 183, 11-19.
3. Lee, C. L., & Brimblecombe, P. (2016). Anthropogenic contributions to global carbonyl sulfide, carbon disulfide and organosulfides fluxes. *Earth-science reviews*, 160, 1-18.

Response to Reviewer #2

We would like to thank reviewer #2 for their time in reviewing this manuscript.

Comments

The uptake of OCS is tied to the OCS concentration within the canopy. There are large variations in OCS uptake as OCS depleted air flows through vegetation, e.g. Berkelhammer et al. 2020 (<https://doi.org/10.1029/2019GL085652>). Most canopies do not see free troposphere concentrations of OCS. This will affect the plant uptake flux component substantially and should be addressed.

This comment was also made by reviewer #1. Because we use an inversion framework, we employ a zero-order flux approach, which is technically easy to implement. Indeed, we find adjustments that point to a large drawdown of COS in the canopy. Like we mention in the discussion, in analyzing the flux adjustment, we should be aware that part of the adjustment is due to the “concentration” effect. Based on the comments of both reviewers, we will make this issue clearer in the revised manuscript.

DMS should not be considered a major source of OCS. Many researchers still refer to the 7% yield figure from the Barnes 1996 paper, but note that chamber studies proceeded without NO_x and at high DMS concentrations. Subsequent studies demonstrated that an alternative chemical pathway is typically taken, and that changes in NO_x affect OCS formation greatly (e.g. [https://doi.org/10.1016/S1352-2310\(98\)00120-4](https://doi.org/10.1016/S1352-2310(98)00120-4)). In other words, if you would like to include DMS, it is important to also model NO_x. This is most likely such a small contribution that in the Whelan et al., 2018 synthesis, it was concluded that DMS should only be included as a source of uncertainty in the ocean flux rather than a source itself.

Currently we do not simulate NO_x chemistry in the COS inverse modelling, since this largely increases the complexity of the modelling system. However, DMS is normally emitted over remote oceans, where low NO_x concentrations prevail. We do not fully agree with the reviewer that the COS yield of 0.7% should not be included as long as the COS budget is not closed. Nevertheless, we will perform a NO-DMS inversion in which we place the DMS-related COS emissions to the “unknown” category.

If the lifetime of CS₂ was 12 days, it might make sense to model CS₂ separately, since the associated OCS will not show up in the air parcel until it has traveled nearly around the globe. However, more recent evidence suggested that the lifetime is much shorter than that. For example, see the 3D atmospheric transport study performed by Anwar Khan which focusses only on CS₂ and estimates a lifetime of less than 4 days at maximum: [10.3934/environsci.2017.3.484](https://doi.org/10.3934/environsci.2017.3.484).

We thank the reviewer to point out this publication. We have checked the rate constant used in our paper for CS₂+OH, which is $2 \times 10^{12} \text{ cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$ (line 214). Assuming that the OH average concentration is about $10^6 \text{ molecules cm}^{-3}$, we recalculated the lifetime as about 5.79 days. We cited the reference Khalil & Rasmussen (1984), but the rate constant is from another paper (Jones et al., 1983). The lifetime quoted in the paper will be corrected to ~6 days accordingly. Khan et al. (2017) presented a global modelling study of CS₂ and quote a lifetime of 2.8-3.4 days, based on a different evaluation of the CS₂+OH rate (Sander et al., 2006). This lifetime is still half of our lifetime, and because reviewer #1 also pointed to the novelty of

including the CS₂ precursor, we will present a sensitivity study for the CS₂ lifetime in the revised manuscript.

The SiB3 model had a known phenology problem, where the growing season starts too soon (by a couple of weeks, perhaps) and ends too early. Does SiB4 have this issue? It would be good to check the seasonal timing by comparing to SiB3 or even another slightly complicating proxy, e.g. SIF.

Our biosphere fluxes are based on simulations with the Simple Biosphere model, version 4 (SiB4) (Haynes et al., 2019). New in SiB4 compared to SiB3 (used by Berry et al. (2013)) are capabilities to simulate i.e. carbon pools, land cover heterogeneity, and leaf phenology. Further, COS uptake formulations in SiB4 are the same as in Berry et al. (2013). One of the new features in SiB4 compared to SiB3 is that it includes prognostic phenology, and this phenology is no longer depending on satellite products. So, if SiB3 had a problem with the phenology this does not automatically mean that SiB4 has the same problem. In fact, from an ongoing comparison with field observations the phenology looks good in SiB4. Results will be presented in a manuscript that is currently in preparation by co-author Linda Kooijmans. Therefore, we consider further discussion of the prior SiB4 fluxes beyond the scope of this study.

The abstract should be revisited to better reflect the conclusions of the study and to tidy up the language, e.g. sources of OCS are obviously included in current budgets.

The abstract will be revised to better show the conclusions and the cohesion of language.

In short, revising the inclusion of DMS and addressing the first order plant uptake issue will certainly create a different budget overall, and may affect the conclusions.

We will discuss these issues in the revised manuscript. We argue, however, that a zero-order approach is sufficient for a first inverse modelling study, since the fluxes are allowed to adjust to the concentration effect. Concerning DMS, we consider this issue currently unresolved, and we will include a NO-DMS inversion.

References:

1. Khalil, M. A. K., & Rasmussen, R. A. (1984). Global sources, lifetimes and mass balances of carbonyl sulfide (OCS) and carbon disulfide (CS₂) in the earth's atmosphere. *Atmospheric Environment* (1967), 18(9), 1805-1813.
2. Jones, B. M. R., Cox, R. A., & Penkett, S. A. (1983). Atmospheric chemistry of carbon disulphide. *Journal of Atmospheric Chemistry*, 1(1), 65-86.
3. Khan, A., Razis, B., Gillespie, S., Percival, C., & Shallcross, D. (2017). Global analysis of carbon disulfide (CS₂) using the 3-D chemistry transport model STOCHEM. *Aims Environ. Sci*, 4, 484-501.
4. Sander SP, Friedl RR, Golden DM, et al. (2006) Chemical kinetics and photochemical data for use in atmospheric studies. Evaluation number 15, JPL publication 06-2, Jet Propulsion Laboratory, Pasadena, CA.
5. Berry, J., Wolf, A., Campbell, J. E., Baker, I., Blake, N., Blake, D., ... & Stimler, K. (2013). A coupled model of the global cycles of carbonyl sulfide and CO₂: A possible new window on the carbon cycle. *Journal of Geophysical Research: Biogeosciences*, 118(2), 842-852.
6. Haynes, K. D., Baker, I. T., Denning, A. S., Stöckli, R., Schaefer, K., Lokupitiya, E. Y., & Haynes, J. M. (2019). Representing grasslands using dynamic prognostic

phenology based on biological growth stages: 1. Implementation in the Simple Biosphere Model (SiB4). *Journal of Advances in Modeling Earth Systems*, 11(12), 4423-4439.

Additional response to reviewer #2

We have an additional short response to the following comment, based on our recent analysis of the CS₂ lifetime (Khan et al. 2017).

Comments

If the lifetime of CS₂ was 12 days, it might make sense to model CS₂ separately, since the associated OCS will not show up in the air parcel until it has traveled nearly around the globe. However, more recent evidence suggested that the lifetime is much shorter than that. For example, see the 3D atmospheric transport study performed by Anwar Khan which focusses only on CS₂ and estimates a lifetime of less than 4 days at maximum: [10.3934/environsci.2017.3.484](https://doi.org/10.3934/environsci.2017.3.484).

We thank the reviewer for pointing out this more recent paper regarding the CS₂ lifetime. We implemented the CS₂ + OH chemistry using the rate quoted in (Khan et al. 2017). As a result, we spotted a mistake in the rate constant of the CS₂ + OH reaction in the Khan et al. (2017) paper. According to (Khan et al. 2017), the rate constant of CS₂+OH is (their Table 1):

$$\text{rate constant 1} = \frac{1.25 \times 10^{-16} e^{\frac{4550}{T}}}{(1 + 1.81 \times 10^{-13} e^{\frac{3400}{T}})}$$

However, the rate constant (Sander et al., 2006, Hynes et al., 1988) should read:

$$\text{rate constant 2} = \frac{1.25 \times 10^{-16} e^{\frac{4550}{T}}}{(T + 1.81 \times 10^{-3} e^{\frac{3400}{T}})}$$

We verified with the authors of Khan et al. (2017) that this is a typo and that the rate constant is correctly implemented in their model. We also implemented the Sander et al. (2006) rate constant in our simulations. Results are shown in Figure 1.

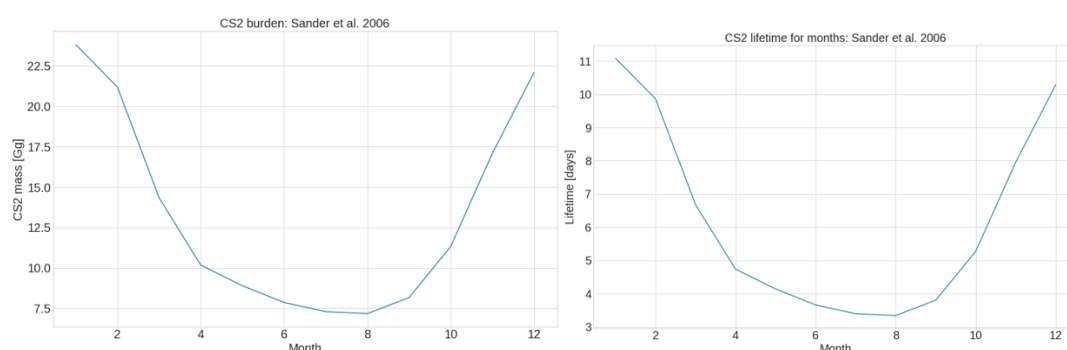


Figure 1: Monthly averaged burden (left) and atmospheric lifetime (right) of CS₂, calculated in TM5 with the rate constant of Sander et al. (2006) implemented.

As can be seen in Figure 1, the burden and lifetime vary considerably over the year. The yearly average lifetime (burden/loss) amounts to 6.2 days, substantially larger than the value quoted in Khan et al. (2017): 2.8-3.4 days. Possible reasons are (1) our burden is higher due to larger

emissions and possibly lower OH (2) no deposition has been implemented in our simulations. In Khan et al. (2017) loss through deposition in their standard run account for ~15% of the CS₂ removal. Note also that their standard model simulation underestimated CS₂ mole fractions, specific over large emission regions.

For comparison, Figure 2 shows the results from our standard run, with a constant CS₂ + OH rate from Jones et al. (1982): $2.0 \times 10^{-12} \text{ cm}^3 \text{ molecules}^{-1} \text{ s}^{-1}$.

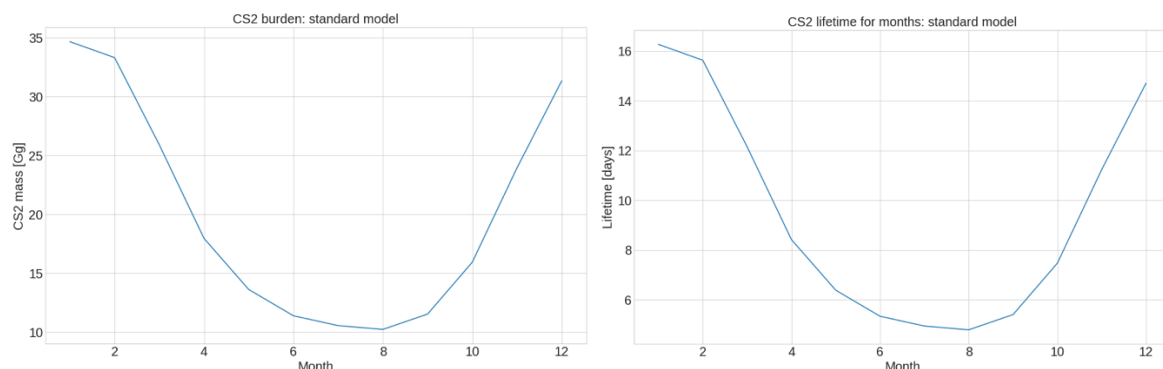


Figure 2: Monthly averaged burden (left) and atmospheric lifetime (right) of CS₂, calculated in TM5 with the rate constant of Jones et al. (1982) implemented.

As can be seen, the lifetime and burden are slightly larger using this rate. The yearly average lifetime amounts to 9.4 days. Figures 3 and 4 show the January and July COS mole fraction difference at the surface between our standard model and the Sander et al. (2006) rate. It can be seen that the differences are relatively small and remain smaller than 10 pmol mol⁻¹.

Since the lifetime in TM5 is relatively long (> 10 days in the Northern Hemisphere winter), we would argue that it is necessary to simulate CS₂ as a separate tracer, to model the delayed COS production from CS₂ oxidation. To highlight this further, we will (in the Supplement of the revised paper) show results from a simulation in which we emit CS₂ directly as COS.

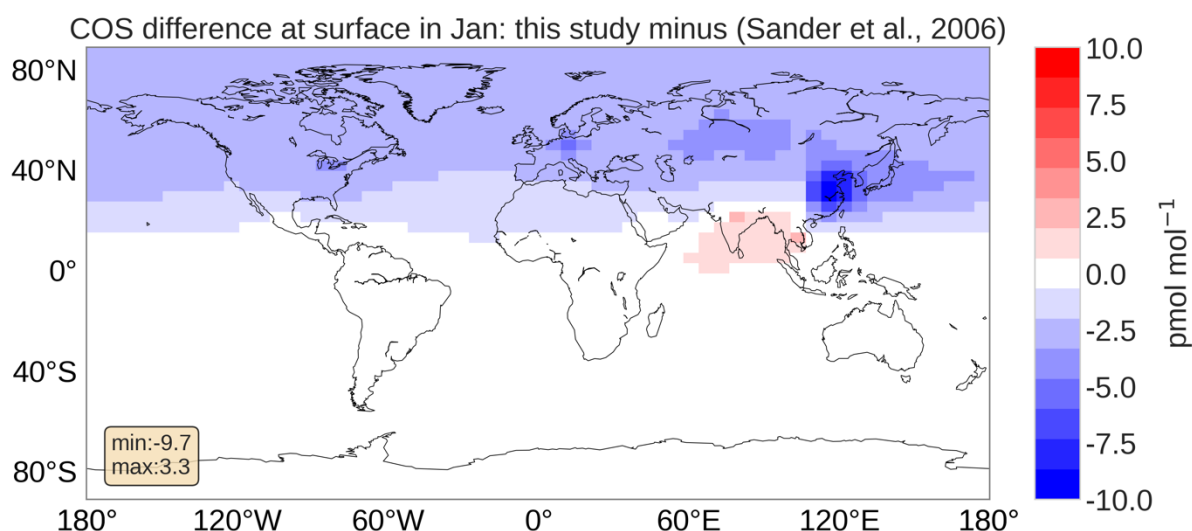


Figure 3. COS mole fractions difference between this study and the rate as (Sander et al., 2006). This is the average COS mole fraction difference in January 2008-2010. The maximum and minimum values of the difference are marked in the left bottom corner.

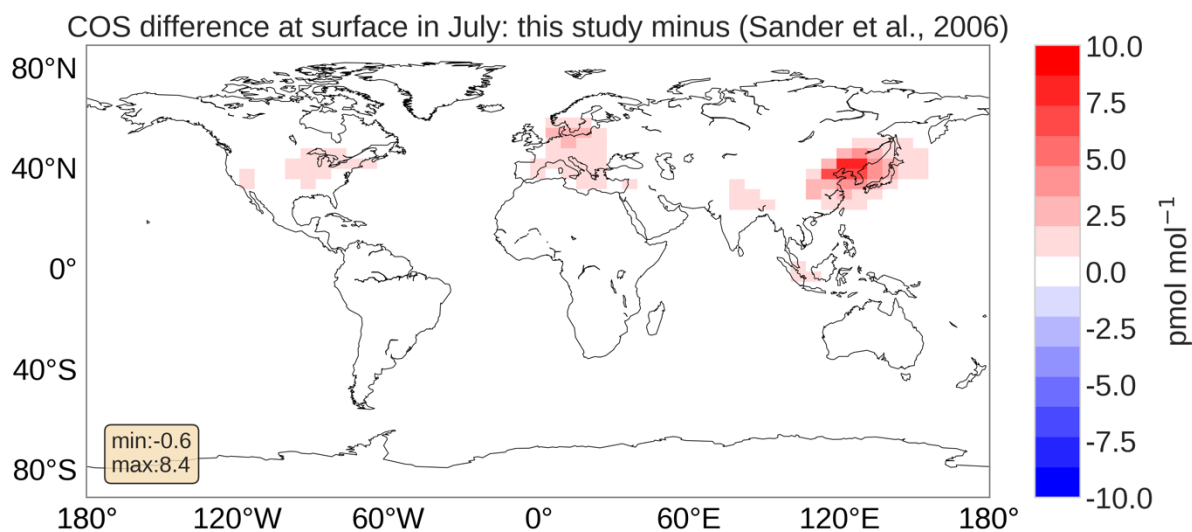


Figure 4. The same as Figure 3 but for July 2008-2010.

Reference:

Khan, Anwar, et al. "Global analysis of carbon disulfide (CS₂) using the 3-D chemistry transport model STOCHEM." *Aims Environ. Sci* 4 (2017): 484-501.

Sander, S. P., et al. Chemical kinetics and photochemical data for use in atmospheric studies evaluation number 15. Pasadena, CA: Jet Propulsion Laboratory, National Aeronautics and Space Administration, 2006.

Hynes, Anthony J., P. H. Wine, and J. M. Nicovich. "Kinetics and mechanism of the reaction of hydroxyl with carbon disulfide under atmospheric conditions." *The Journal of Physical Chemistry* 92.13 (1988): 3846-3852.

Khalil, M. A. K., and R. A. Rasmussen. "Global sources, lifetimes and mass balances of carbonyl sulfide (OCS) and carbon disulfide (CS₂) in the earth's atmosphere." *Atmospheric Environment* (1967) 18.9 (1984): 1805-1813.

Jones, B. M. R., et al. "OCS formation in the reaction of OH with CS₂." *Chemical Physics Letters* 88.4 (1982): 372-376.

Inverse modelling of carbonyl sulfide: implementation, evaluation and implications for the global budget

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Abstract.

Carbonyl sulfide (COS) has the potential to be used as a climate diagnostic due to its close coupling to the biospheric uptake of CO₂ and its role in the formation of stratospheric aerosol. The current understanding of the COS budget, however, lacks COS sources, which have previously been allocated to the tropical ocean. This paper presents a first attempt of global inverse modelling of COS within the 4-Dimensional variational data-assimilation system of the TM5 chemistry transport model (TM5-4DVAR) and a comparison of the results with ~~independent-various~~ COS observations. We focus on the global COS budget, including COS production from its precursors carbon disulfide (CS₂) and dimethyl sulfide (DMS). To this end, we implemented COS uptake by soil and vegetation from an updated biosphere model (SiB4), ~~and~~. In the calculation of these fluxes, a fixed atmospheric mole fraction of 500 pmol mol⁻¹ was assumed. We also used new inventories for anthropogenic and biomass burning emissions. The model framework is capable of closing the COS budget by optimizing for missing emissions using NOAA observations in the period 2000–2012. The addition of 432 Gg S a⁻¹ COS is required to obtain a good fit with NOAA observations. This missing source shows little year-to-year variations, but considerable seasonal variations. We found that the missing sources are likely located in the tropical regions, and an overestimated biospheric sink in the tropics cannot be ruled out due to missing observations in the tropical continental boundary layer. Moreover, high latitudes in the Northern Hemisphere require extra COS uptake or reduced emissions. HIPPO aircraft observations, NOAA airborne profiles from an ongoing monitoring program, and several satellite data sources are used to evaluate the optimized model results. This evaluation indicates that COS mole fractions in the free troposphere ~~remains-remain~~ underestimated after optimization. Assimilation of HIPPO observations slightly improves this model bias, which implies that additional observations are urgently required to constrain sources and sinks of COS. We finally find that the biosphere flux dependency on surface COS mole fraction (which was not accounted for in this study) may substantially lower the fluxes of the SiB4 biosphere model over strong uptake regions. Using COS mole fractions from our inversion, the prior biosphere flux reduces from 1053 Gg S a⁻¹ to 851 Gg S a⁻¹, which is closer to 738 Gg S a⁻¹ as was found by Berry et al. (2013). In planned further studies we will implement this biosphere

dependency, and additionally assimilate satellite data with the aim to better separate the role of the oceans and the biosphere in the global COS budget.

25 1 Introduction

Carbonyl sulfide (COS or OCS) is a low abundant trace gas in the atmosphere with a lifetime of about 2 years and a tropospheric mole fraction of about 484 pmol mol⁻¹ (Montzka et al., 2007). COS is regarded as a promising diagnostic tool for constraining photosynthetic gross primary production (GPP) of CO₂ through similarities in their stomatal control (Montzka et al., 2007; Campbell et al., 2017; Berry et al., 2013; Whelan et al., 2018; Kooijmans et al., 2017, 2019; Wang et al., 2016). COS also contributes to stratospheric sulfur aerosols, which have a cooling effect on climate and hence mitigate climate warming (Crutzen, 1976; Andreae and Crutzen, 1997; Brühl et al., 2012; Kremser et al., 2016). In the recent decades, COS mole fractions in the troposphere have remained relatively constant, which implies that sources and sinks of COS are balanced. Whelan et al. (2018) reviewed the state of current understanding of the global COS budget and the applications of COS to ecosystem studies of the carbon cycle. The most pressing challenge currently is to ~~close the global COS budget~~reconcile the balance of COS sources and sinks, given the small global atmospheric trends.

Previous studies show that substantial emissions of COS are coming from oceanic, anthropogenic, and biomass burning sources, and the largest sinks are uptake by plants and soils (Watts, 2000; Kettle et al., 2002; Berry et al., 2013). Oceanic emissions are thought to be the largest source of COS, both directly and indirectly, due to emissions of CS₂ and ~~DMS (Lennartz et al., 2017, 2019)~~possibly DMS (Lennartz et al., 2017, 2020), which can be quickly oxidized to COS in the atmosphere (Sze and Ko, 1980). There are considerable uncertainties related to this indirect COS source, with reported yields to COS being (83 ± 8)% from CS₂ (Stickel et al., 1993), and (0.7 ± 0.2)% from DMS under ~~low-NO_x~~NO_x-free conditions at 298 K (Barnes et al., 1996). Blake et al. (2004) reported anthropogenic Asian emissions for COS and CS₂, which appear to have been underestimated by 30–100% due to underestimated coal burning in China (Du et al., 2016). Zumkehr et al. (2018) recently presented a new global anthropogenic emission inventory for COS. The new anthropogenic emission estimates are, with 406 Gg S a⁻¹ in 2012, substantially larger than the previous estimate of 180.5 Gg S a⁻¹ by (Berry et al., 2013). Another recent study (Stinecipher et al., 2019) concluded that it is unlikely that biomass burning accounts for the balance between sources and sinks of COS, due to the relatively small contribution of biomass burning to the total emissions ((60±37) Gg S a⁻¹).

Suntharalingam et al. (2008) made a first attempt to simulate the global COS budget using the GEOS-Chem model and global-scale surface measurements from NOAA. In order to fit the observed seasonal cycle of COS mole fraction, they had to double the terrestrial vegetation uptake estimated in (Kettle et al., 2002), reduce the southern extra-tropical ocean source, and assume an additional COS source of 235 Gg S a⁻¹. Campbell et al. (2008) found that this upward revision could be validated using direct observations from the continental boundary layer from the intensive INTEx-NA airborne campaign. Berry et al. (2013) implemented COS in the global biosphere model SiB3. They inferred that, in order to compensate for updated COS biosphere and soil sinks of 1093 Gg S a⁻¹, there must be additional COS sources of 600 Gg S a⁻¹, which was allocated to the

ocean. Glatthor et al. (2015) and Launois et al. (2015) estimated direct COS emissions from the ocean as 992 and 813 Gg S a⁻¹, respectively, and also Kuai et al. (2015) hinted to underestimated COS sources from tropical oceans by optimizing sources using one month of COS satellite observations by TES-Aura. However, [Lennartz et al. \(2017\)](#) [Lennartz et al. \(2017, 2019\)](#) used COS measurements in ocean water to show that the direct oceanic emissions were much lower (130 Gg S a⁻¹) than top-down studies suggested. It is therefore not resolved whether ocean emissions account for the missing source.

In this paper, we address several important open questions concerning the COS budget using inverse modelling techniques, employing the TM5-4DVAR modelling system. We focus on the closure of the COS budget, the contributions of the potential COS precursors CS₂ and DMS, and evaluation of the results with aircraft and satellite observations. In Section 2 we will describe the observations, the implementation of COS, CS₂, and DMS in TM5, and the inverse modelling system TM5-4DVAR. In Section 3, we will analyze the results of various inverse model calculations, which are discussed further in Section 4.

2 Method

This study aims to close the [gap in the](#) global COS budget by so-called flux inversions. This technique employs atmospheric measurements to optimize sources and sinks of trace gases such that mismatches between simulations and observations are minimized. In Section 2.1 the observations used in this study are introduced. Section 2.2 will subsequently describe our modelling system, including new emission data sets that have been coupled to the modelling system. The inverse modelling framework is discussed in Section 2.3.

2.1 Measurements

2.1.1 NOAA flask data

The NOAA surface flask network provides long-term measurements of COS mole fraction at 14 locations at weekly–monthly frequencies. Most of the stations are located in the Northern Hemisphere (NH), as shown in Figure 1. Although the number of sampling sites is modest, their locations cover most latitudinal regions, and sample over both land and coastal area. [It is worth to note that there is a lack of observations in the tropical continental boundary layer.](#) The observational error for each [site sample](#) is relatively small (< 7 pmol mol⁻¹), therefore we have taken inter-annual variability of COS from Table 1 in Montzka et al. (2007) to represent a fixed observational error upper-limit at each site. In general, the observational error defined in this way varies between 4–10 pmol mol⁻¹ in the NH, and between 2–4 pmol mol⁻¹ in the Southern Hemisphere (SH). This error is used in the inverse modelling as will be described in Section 2.4.

2.1.2 HIPPO aircraft and NOAA airborne data

Flask data of the HIAPER Pole-to-Pole Observations (HIPPO) experiments (Wofsy, 2011; Wofsy et al., 2017) is used to validate the results of the inverse modelling. There are five HIPPO campaigns conducted from 2009 to 2011 that sampled the

COS mole fraction from the North Pole to the South Pole, and from the lower troposphere up to the stratosphere. Three different instruments were used to make measurements of COS during HIPPO. Instrument 2 was used by NOAA to measure COS, and instrument 1 was calibrated consistently with the NOAA calibration standard. ~~Instrument~~ Results of instrument 3 is an independent instrument, but the were scaled to be consistent with that of instrument 2, such that results from all three instruments on HIPPO are referenced to the same NOAA scale. The probability distribution function of the mole fractions ~~shows~~ confirms that the three instruments ~~are measuring report~~ consistent values, with similar averages (see Figure S1). Thus, HIPPO data provides valuable data to check the consistency of the optimized COS budget. The flight routes of the five campaigns are shown in Figure 1. In some numerical experiments, HIPPO data are additionally assimilated to investigate their impact on the optimized COS budget. To investigate this impact on the vertical distribution of COS, we compared to 2008–
 90 2011 NOAA airborne data that are mainly available over North America (Figure 1). The number of aircraft sites used is 19, and the upper altitude that was typically reached is 8 km.

2.1.3 Satellite data

Our inverse modelling results are compared to three independent satellite data sources: TES-Aura, ACE-FTS, and MIPAS. We have selected the period 2008–2011 for the comparison.

100 NASA’s Tropospheric Emission Spectrometer (TES) is a both nadir and limb viewing instrument that flies on the AURA satellite, which was launched in 2004 (Beer et al., 2001). TES measures the infrared radiation emitted from the Earth and atmosphere in high spectral resolution for 16 orbits every other day. From these spectra, abundances of tropospheric trace gases are retrieved. The COS product used in this study is described in Kuai et al. (2014). The COS retrievals cover the whole vertical column and have less than 1 degree of freedom (DOF) and show maximum sensitivity in the 300–500 hPa region. We
 105 will therefore focus our comparisons on total COS columns. To account for the non-uniform vertical sensitivity, we use the Averaging Kernel (AK) in the model–satellite comparison. As described in Kuai et al. (2014), the AK included in the TES data files is defined in log-space, and should be applied as:

$$\ln(\chi_{con}) = \ln(\chi_p) + \mathbf{A}[\ln(\chi_m) - \ln(\chi_p)], \quad (1)$$

where χ_{con} , χ_p , and χ_m are respectively the convolved, prior, and modeled profiles, and \mathbf{A} is the AK. In Section 3.4 the
 110 modeled profiles are convolved with the TES AK (χ_{con}), vertically integrated, and compared to the TES columns.

The Atmospheric Chemistry Experiment Fourier Transform Spectrometer (ACE-FTS) is a high spectral resolution infrared FTS instrument that performs solar occultation measurements, with the aim to sample stratospheric and upper tropospheric profiles of trace gases (Boone et al., 2013). The instrument flies on SCISAT, a Canadian satellite mission for remote sensing of the Earth’s atmosphere that was launched in 2003. Its orbit covers tropical, mid-latitude, and polar regions. COS is one of
 115 the atmospheric trace gases measured by the ACE-FTS instrument (Koo et al., 2017). ACE-FTS profiles have been compared to balloon observations and generally showed good agreement, with underestimations smaller than 20% (Krysztofiak et al., 2015). We use product version 3.6 and only observations with quality flag of zero are used. ACE-FTS measures COS within 0–150 km vertically, but the data quality is only sufficient in the upper troposphere and lower stratosphere (UTLS).

120 The Michelson Interferometer for Passive Atmospheric Sounding (MIPAS) is a Fourier transform spectrometer for ~~the~~
~~detection of limb emission spectra in~~ detection of the radiative emission of various molecules in limb observation mode in the
middle and upper atmosphere (Fischer et al., 2008). MIPAS flew on ESA’s Envisat platform that operated between 2002–2012.
MIPAS delivers global atmospheric COS profiles in the upper troposphere and stratosphere (Glatthor et al., 2015, 2017). Similar
to TES, the MIPAS ~~product provides data~~ product contains representative AKs and prior profiles to facilitate ~~comparisons~~
comparison to modelled profiles, but not in log-space (~~Stiller et al., 2012~~), since MIPAS COS is evaluated by a linear retrieval:
125

$$\chi_{con} = \chi_p + \mathbf{A}[\chi_m - \chi_p], \quad (2)$$

where χ_m has to be resampled on the MIPAS retrieval grid in advance.

As for most other gases, the prior profile for MIPAS COS retrievals is a zero profile. Eq .2 thus becomes a simple multipli-
cation of the AK with the modelled profiles. A detailed description of the application of MIPAS AKs on other datasets can be
130 found in Stiller et al. (2012).

The MIPAS product has been compared to modelled COS distributions (Glatthor et al., 2015) and ACE-FTS (Glatthor et al.,
2017). The latter comparison showed that MIPAS retrieves higher mole fractions around the tropopause compared to ACE-
FTS. The MIPAS product has also been compared to airborne measurements of the HIPPO, ARCTAS and INTEx-B campaigns
(Supplement of Glatthor et al. 2015). ~~Further~~ Finally, MIPAS has been compared to MkIV and SPIRALE profiles (Glatthor
135 et al., 2017). ~~Comparable to the ACE-FTS product, MIPAS COS profiles are only valid in the UTLS.~~

The retrievals of TES, MIPAS and ACE-FTS v3.6 are provided on 14, 60, and 150 vertical levels in the atmosphere, respec-
tively. We map our modelled COS profiles to these levels using a mass conserving interpolation scheme.

2.1.4 Seasonal decomposition

In Section 3.1 we apply a simple seasonal decomposition method to our calculated exchange fluxes. The seasonal decomposi-
140 tion is performed using Python module StatsModels version 0.10. The time series are decomposed into trend, seasonality and
noise:

$$\mathbf{y}(t) = \mathbf{y}_t(t) + \mathbf{y}_s(t) + \mathbf{y}_r(t), \quad (3)$$

with $\mathbf{y}(t)$ the monthly exchange fluxes, and \mathbf{y}_t , \mathbf{y}_s , and \mathbf{y}_r the trend, seasonal, and residual components, respectively.

2.2 Model description

145 2.2.1 Anthropogenic emissions

We have implemented the anthropogenic emissions based on a recent global gridded emission inventory of COS (~~Zumkehr et al., 2018~~)
(Campbell et al., 2015; Zumkehr et al., 2018). Since we aim to model COS, CS₂ and DMS as separated tracers, we disentan-
gled the reported COS emissions into COS and CS₂ contributions. Here, we applied an assumed yield of 0.87 (Zumkehr et al.,

2018), which means that 1 mole CS₂ yields 0.87 mole COS. As a precursor of COS, CS₂ reacts with OH to produce COS, and
 150 has an atmospheric lifetime of about 12 days (Khalil and Rasmussen, 1984). We applied a detailed anthropogenic emission budget for COS and CS₂ from Table 1 in (Lee and Brimblecombe, 2016). This allows us to roughly estimate the ratio of this budget and hence the direct and indirect COS anthropogenic emissions. The converted emissions averaged over the period 2000–2012 are summarized in Table 1.

The total anthropogenic COS emissions are on average 343.3 Gg S a⁻¹, split between direct COS emissions of 147.5 Gg S
 155 a⁻¹ and CS₂ emissions of 450.2 Gg S a⁻¹. This indicates that CS₂ is an important precursor of COS. Figure 2 shows time-series of COS and CS₂ anthropogenic emissions. COS emissions are dominated by industrial and residential coal sources, while CS₂ emissions are dominated by rayon industry and TiO₂ production. Moreover, while COS emissions remained relatively constant in the 2007–2012 period, CS₂ emissions show an increasing trend.

While Zumkehr et al. (2018) assumed a molar yield of CS₂ to COS of 87%, other reported yields are (83 ± 8)% (Stickel
 160 et al., 1993) and 81% (Chin and Davis, 1993). We decided to use a yield of 83% in our modelling, while we used the reported yield of 87% to produce the numbers listed in Table 1. This implies that we introduce slightly less COS in the atmosphere compared to using the Zumkehr et al. (2018) data as direct COS emissions. Note that we apply all categorical emissions or fluxes with a monthly time resolution. It is also worth noting that the uncertainty in the anthropogenic inventory is much larger than the uncertainty in molar yield.

165 2.2.2 Biomass burning emissions

We estimated biomass burning emissions based on the widely used GFED V.4.1 data set (Randerson et al., 2018) for six of the seven emissions categories listed in Table 2. In converting dry mass burned to COS emissions, we used the updated emission factors reported in Andreae (2019). For biofuel use, we base our estimates on the Community Emission Data System (CEDS) (Hoesly et al., 2018). We calculated COS emissions by first converting CO emissions to dry mass burned, which were converted
 170 to COS emissions in a second step. Emission factors are listed in Table 2. In this process we made a distinction between biofuel with and without dung. Dung burning is mainly employed in South Asia (Fernandes et al., 2007) and we applied the dung emission ratios only in the region 60–100°E and 0–40°N. Our biomass burning emissions in the 2000–2012 period are in the range 118–154 Gg S a⁻¹ (Figure 2), similar to the emissions used in ~~Berry et al. (2013)~~ (Berry et al., 2013) (135 Gg S a⁻¹) ~~–The and estimates reported in (Campbell et al., 2015) (116±52 Gg S a⁻¹).~~ The more recent biomass burning estimate from (Stinecipher et al., 2019) based on GFED 1997–2016 data reports global emissions of 60±37 Gg S a⁻¹. Note, however, that biofuel use is not included in this estimate. The spatial and seasonal distribution of the biomass burning emissions averaged over the period 2000–2012 is presented in ~~supplementary~~ Supplementary Figure S2.

2.2.3 Biosphere flux

Our biosphere fluxes are based on simulations with the Simple Biosphere model, version 4 (SiB4) ~~(Haynes et al., 2019)~~ –SiB4 calculates the uptake by vegetation based on (Berry et al., 2013)–(Berry et al., 2013; Haynes et al., 2019). Currently, soil uptake is scaled to the CO₂ soil respiration term, and the implementation of specific COS soil models (Sun et al., 2015; Ogée

et al., 2016) is ongoing. The SiB4 model was constrained by a prescribed COS mole fraction of 500 pmol mol⁻¹ outside of canopy. ~~This 500 pmol mol⁻¹ is merely as a placeholder, and probably leads to too large fluxes over active biosphere, where COS mole fractions decline because of strong uptake. This is further discussed in Section 3.5.~~ Meteorological data that are used as forcing for the SiB4 model are taken from the Modern Era Retrospective Analysis for Research and Applications (MERRA) and are available from 1980 onwards (Rienecker et al., 2011). A spin-up of the model was performed for the period 1850-1979 to reach an equilibrium of the carbon pools. As no MERRA data were available for the spin-up period, the climatological average of MERRA data over the period 1980-2018 was used as meteorological input for the spin-up period. A final simulation was performed for 1980–2018 with the actual MERRA driver data. The 2000–2018 average flux to the biosphere (vegetation plus soil) amounts to -1053 Gg S a⁻¹, in line with estimates using SiB3 (-951 Gg S a⁻¹ (Kuai et al., 2015)) and (Berry et al., 2013). The spatial and seasonal distribution of the biosphere uptake is shown in Figure S3. The uptake shows a large seasonal cycle in the NH and large uptake over tropical forests. The biosphere fluxes were deployed on a monthly timescale.

2.2.4 Ocean emissions

Climatological ocean emissions of COS and the COS precursors CS₂ and DMS are based on Suntharalingam et al. (2008) and Kettle et al. (2002). Large quantities of COS, DMS, and CS₂ are emitted from open oceans. The estimated DMS emissions are about 22 Tg S a⁻¹, and we note that even if the COS yield from oxidation of DMS is as small as 0.7 % (Barnes et al., 1996), already 156 Gg S a⁻¹ COS is formed. The CS₂ direct emission from oceans is roughly 195 Gg S a⁻¹, yielding 81 Gg S a⁻¹ of COS. When the ocean water is cold enough, oceans can turn into a sink of COS instead of a source (Lennartz et al., 2017). Supplementary Figure S4 shows the spatial distribution of the January and July direct and indirect ocean emissions of COS. Note that our estimates of all COS oceanic emissions as 277 Gg S a⁻¹ are substantially smaller than the estimates of 813 Gg S a⁻¹ by Launois et al. (2015).

2.3 TM5-4DVAR inverse modelling system

We have implemented three tracers (COS, CS₂, and DMS) in the inverse modelling framework TM5-4DVAR (Krol et al., 2005, 2008; Meirink et al., 2008). In brief, the TM5 model is used to convert fluxes, collected in state vector \mathbf{x} , to observations \mathbf{y} :

$$\mathbf{y} = \mathbf{H}(\mathbf{x}), \quad (4)$$

where \mathbf{H} represents the global chemistry transport model TM5. Since the relation between fluxes and observations is currently linear, $\mathbf{y} = \mathbf{H}(\mathbf{x})$ can be written as $\mathbf{y} = \mathbf{H}\mathbf{x}$. In a flux inversion a cost function is minimized. The cost function has the form:

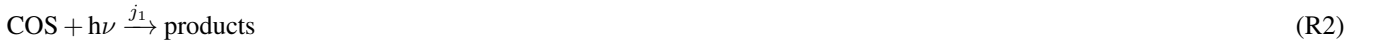
$$J(\mathbf{x}) = \frac{1}{2}(\mathbf{x} - \mathbf{x}_b)^T \mathbf{B}^{-1}(\mathbf{x} - \mathbf{x}_b) + \frac{1}{2}(\mathbf{y} - \mathbf{H}\mathbf{x})^T \mathbf{R}^{-1}(\mathbf{y} - \mathbf{H}\mathbf{x}), \quad (5)$$

where \mathbf{x}_b represents the prior state of the fluxes, and \mathbf{B} and \mathbf{R} are the error covariance matrices of the fluxes and observations, respectively. \mathbf{B} contains the errors assigned to the fluxes, as well as their correlations in space and time (i.e. \mathbf{B} is a non-diagonal

matrix). \mathbf{R} contains the errors assigned to $(\mathbf{y} - \mathbf{H}\mathbf{x})$. These errors are assumed to be uncorrelated and they include, next to the observational errors, also errors related to the process of mapping coarse-scale fluxes \mathbf{x} to localized observations \mathbf{y} . The
 215 adjoint of the TM5 model (Krol et al., 2008; Meirink et al., 2008) is used to calculate the gradient of this cost function with respect to all elements in the state vector:

$$\nabla J(\mathbf{x}) = \mathbf{B}^{-1}(\mathbf{x} - \mathbf{x}_b) + \mathbf{H}^T \mathbf{R}^{-1}(\mathbf{H}\mathbf{x} - \mathbf{y}). \quad (6)$$

In all inversions, \mathbf{y} is represented by COS observations from the NOAA flask network data (Montzka et al., 2007). Our flux space, however, in addition to COS emissions, may represent CS₂ and DMS emissions from anthropogenic activity and oceans.
 220 To map their influence on simulated COS observations \mathbf{y} , we need to consider chemical conversions of CS₂ and DMS to COS. CS₂ and DMS are short-lived trace gases, with atmospheric lifetimes of approximately 12 days (Khalil and Rasmussen, 1984) and 1.2 days (Khan et al., 2016; Boucher et al., 2003; Breider et al., 2010), respectively. For CS₂ we implemented OH-initiated conversion to COS, while for DMS we simply apply exponential decay with a lifetime of 1.2 days. COS itself is also destroyed by OH in the troposphere and by photolysis in the stratosphere. For OH, we use monthly varying climatological OH fields
 225 (Spivakovsky et al., 2000), and applied a correction factor of 0.92 (Naus et al., 2019). In summary, the chemistry that is implemented therefore consists of the following four reactions:



230



where j_1 is the stratospheric photolysis frequency, and r_1 and r_2 are the rate constants of COS and CS₂ OH-oxidation, respec-
 235 tively. The fractions f_1 and f_2 represent the molar yields of COS from CS₂ (taken as 0.83 (Stickel et al., 1993)) and DMS (taken as 0.007 (Barnes et al., 1996)). The rate r_1 is calculated according to the Arrhenius equation:

$$r_1 = A e^{\frac{-1200 \text{ K}}{T}}, \quad (7)$$

where T is temperature in K, and A is $1.13 \times 10^{-12} \text{ cm}^3 \text{ s}^{-1} \text{ molecule}^{-1}$ (Cheng and Lee, 1986). The rate r_2 is $2.0 \times 10^{-12} \text{ cm}^3 \text{ s}^{-1} \text{ molecule}^{-1}$ (Jones et al., 1983). Note that this rate expression is independent of temperature and slightly

240 different from Sander et al. (2006). This latter rate was used in Khan et al. (2017), and resulted in a short CS₂ lifetime of 2.8–3.4
days. However, when we implement the Sander et al. (2006) are in TM5, we find a CS₂ lifetime of 6.2 days. This might be
due to the fact that we ignore CS₂ deposition (15% of the loss according to Khan et al. (2017)) or that we use lower OH or
higher emissions. Rate $r_2 = 2.0 \times 10^{-12} \text{ cm}^3 \text{ s}^{-1} \text{ molecule}^{-1}$ leads to an atmospheric CS₂ lifetime of 9.4 days in TM5. Rate
 r_3 represents an exponential decay of 1.2 days for DMS ($r_3 = 9.6 \times 10^{-6} \text{ s}^{-1}$).

245 COS photolysis frequencies are calculated based on troposphere ultraviolet and visible (TUV) radiation model (Madronich
 et al., 2003). Based on monthly climatologies of ozone profiles and temperatures, monthly-averaged photolysis frequencies are
 calculated on a 1 km grid spanning 0 – 120 km and on 180 latitude bands. Implemented in TM5, COS loss in the stratosphere
 amounts to about 40 Gg S a⁻¹. This estimate is in line with earlier estimates of (50±15) Gg S a⁻¹ (Brühl et al., 2012; Barkley
 et al., 2008; Chin and Davis, 1995; Engel and Schmidt, 1994; Weisenstein, 1997; Krysztofiak et al., 2015; Turco et al., 1980;
 250 Crutzen and Schmailzl, 1983; Crutzen, 1976).

2.4 Model-data mismatch errors

The diagonal elements of the error covariance matrix **R** in Eq. 5 contain contributions from observational errors, representation
 errors and errors related to applying large fluxes in the planetary boundary layer (Bergamaschi et al., 2010):

$$\sigma_t = \sqrt{(\sigma_o^2 + \sigma_r^2 + \sigma_f^2)}, \quad (8)$$

255 where σ_t is the total error, σ_o the observational error, σ_r the representation error, and σ_f an error related to applying large
 surface fluxes. The assumed observational error is shown in Figure 3. It is worth to note that observational errors are usually
 overwhelmed by the representation and flux errors. The representation error is calculated by sampling the modelled gradients
 in the vicinity of the sampled station (Bergamaschi et al., 2010). Finally, the flux error in each cell is linked to the magnitude
 of the monthly surface flux f (kg m⁻² s⁻¹ in each cell) applied in the model as:

$$260 \quad \sigma_f = \frac{f g M_{\text{air}} \Delta t}{M_S \Delta p}. \quad (9)$$

Here, f represent the sum of all COS prior flux components. In this sum, the biosphere flux is dominant over regions with
 strong biosphere uptake. Further, g is gravitational acceleration (9.8 m s⁻²), M_{air} is molar mass of dry air (28.9 kg kmol⁻¹),
 M_S is molar mass of sulfur (32.1 kg kmol⁻¹), Δp (Pa) is the thickness of the first model layer, and Δt is the time (s) over
 which the COS flux accumulates (we use 1 hour). Note that σ_f is unit-less and is multiplied by 1×10^{12} to obtain units of pmol
 265 mol⁻¹.

Based on the total error, we define a χ^2 metric to quantify how well the observations are reproduced by the model (e.g. at a
 particular station).

$$\chi^2 = \frac{\sum_{i=1}^N (\mathbf{H}\mathbf{x} - \mathbf{y})^2}{N\sigma_t^2}, \quad (10)$$

where N is the number of individual observations. We can calculate this metric before optimization (prior) and after opti-
 270 mization (posterior). χ^2 is used to diagnose whether inversions are over-fitting or under-fitting the information contained in

the measurement network. A value $\chi^2 \approx 1.0$ indicates that the inverse system is able to fit the data within the error setting (Hooghiemstra et al., 2011). A large posterior χ^2 indicates that the state does not have enough degrees of freedom to fit the observations properly (or the error settings are too small). A small posterior χ^2 indicates over-fitting of the observations (or too wide error settings).

275 2.5 Model settings

In this study, the TM5-4DVAR system is employed on a global resolution of $6^\circ \times 4^\circ$ (longitude \times latitude). Flux fields are coarsened from a resolution of $1^\circ \times 1^\circ$. To create a reasonable start field for the inversions, we initially performed an 11-year forward simulation starting with zero initial mole fractions and baseline surface fluxes augmented by 432 Gg S a^{-1} , distributed uniformly to close [a gap in](#) the global budget. After 11 years, sources and sinks are roughly in balance, with atmospheric mole
280 fractions of about $500 \text{ pmol mol}^{-1}$. Note that fluxes are used as zero-order terms, while the COS removal by OH and photolysis are first order removal terms that grow as the atmospheric COS increases.

We will present the results of four inversions. Firstly, we optimized the missing emissions, which amount to 432 Gg S a^{-1} . This inversion will be denoted by S_u throughout the paper. The aim of this inversion is to investigate the spatial structure and temporal variability of the missing COS emissions. This is the first time that a formal 4DVAR approach is applied to
285 the COS budget. To this end, we start from an emission field of 432 Gg S a^{-1} that is uniformly distributed globally. We optimize emissions on a monthly timescale, and assign a grid-scale prior error of 100%, which is an arbitrary number to give fluxes enough freedom to adjust. In a three-year inversion, the total number of state vector elements amounts to 97200 (36 months \times 45 latitudinal bins \times 60 longitudinal bins). The total number of NOAA observations is much smaller, thus rendering the inversion under-determined. We therefore also use inversion S_u to explore different settings of the temporal and spatial
290 correlation lengths, which control the degrees of freedom of the state vector. We explore spatial correlation lengths of 1000 km, 4000 km, 6000 km, 10000 km, and 20000 km, and temporal correlation lengths of 5.5, 7, 9.5, 12 months.

Secondly, we explore the optimization of specific categories in inversions S1–S3. In S1 we attempt to perform an "objective" inversion, in which we assign grid-scale errors of 50% to the biosphere and ocean (we optimize both COS and CS_2), and 10% to the anthropogenic COS and CS_2 emissions, and to the biomass burning emissions. Furthermore, in S2 we only optimise ocean
295 exchange, and in S3 we only optimise the biosphere exchange. The aim of inversions S1–S3 is to explore whether either ocean fluxes or the biosphere fluxes (or both) should be used to close the [gap in the](#) COS budget. Note that DMS ocean emissions are not optimized. The names and setting of the inversions are summarized in Table 3.

The cost function is minimized with Congrad, an efficient numerical algorithm for solving linear systems (Lanczos, 1950). This minimizer was also used in previous inverse modelling studies with the TM5-4DVAR system (Basu et al., 2013; Monteil
300 et al., 2011, 2013; Houweling et al., 2014; Pandey et al., 2015). For convergence, we request a gradient norm reduction of 1×10^5 , and this reduction is usually achieved within 40 iterations.

We perform flux inversions for the period 2000–2012. To decrease computational costs, we adopt the strategy to run parallel 3-year inversions, and we discard the optimized fluxes of the first 6 months (spin-up) and the last 6 months (spin-down). For example, the first inversion targets the period 1-1-2000 to 1-1-2003, the second inversion 1-1-2002 to 1-1-2005, and so on. In

305 the spin-up period the fluxes in the first 6 months are used to adjust the imperfect initial condition. In the spin-down period, fluxes are less reliable, because they have not been well constrained by observations. The optimized fluxes in the overlapping years are used to check the inversion results for consistency. In general, it is found that the optimized fluxes in the overlapping periods are highly consistent.

3 Results

310 3.1 Closing a gap in the COS budget

In this section, we consider inversion S_u , in which a uniform field emitting 432 Gg S a^{-1} is optimized. We use different settings for the spatial and temporal correlation lengths of this field in the inversion, and quantify the posterior goodness-of-fit using the χ^2 metric (Eq. 10). As presented in more detail in Supplementary Figure S5 we find, as expected, that χ^2 decreases with increasing degrees of freedom (smaller correlations).

315 Overall, the posterior fit to NOAA ~~observations~~ surface observations from 14 sites does not improve significantly for smaller correlation lengths. If we analyse the posterior fit to ~~independent data~~ the short-term sampling program from HIPPO, however, we find that the χ^2 reaches a minimum (see supplement Figure S5). After this minimum, χ^2 values increase again, a possible sign of over-fitting. We therefore select 4000 km and 12 months for respectively the spatial and temporal correlation length, and use these values in the remainder of this study.

320 Figure 4 presents the fit to observations of the prior and posterior simulation, for the inversion with temporal and spatial correlation lengths of 12 months and 4000 km, respectively. Corresponding χ^2 metrics per station are listed as labels in Figure 4. Posterior fits are by design much better than prior fits. Only for NOAA stations THD and NWR the posterior χ^2 remains larger than 3, indicating insufficient degrees of freedom to resolve remaining discrepancies, underestimated model errors, or the influence of outliers (see Figure 4 g, h). THD is a coastal site (107 m a.s.l.) and NWR is a tundra site above
325 treeline (3526 m a.s.l.) in the US (Figure 1), and thus the model resolution of $6^\circ \times 4^\circ$ ~~might be~~ is likely too coarse to represent these sites. ~~Indeed, observations regularly exhibit very low mole fractions during the growing seasons~~ The local coastal effect might be another reason why THD yields a larger χ^2 (Riley et al., 2005). It is worth to note that the posterior simulation does not exhibit jumps in overlapping years from the parallel running inversions, indicating that our inversion strategy works well.

The correlation settings have a large impact on the optimized fluxes. Figure 5 shows the spatial distribution of the posterior
330 flux field calculated with two different correlation settings. For correlations of 1000 km and 5.5 months (panel (a)) we detect a typical pattern that signals over-fitting of the observations. In such a pattern, the optimized flux displays hot spots close to measurement locations (e.g. THD, MLO, SMO). For very long spatial correlations, e.g., 20000 km, posterior fits are poor ($\chi^2 > 6$, see Figure S5) and optimized flux patterns show irregular behaviour (Figure S6). Our best inversion (4000 km and 12 months) produces a smooth optimized flux without apparent spatial patterns near observational stations (Figure 5b). This
335 pattern confirms the missing COS sources in the tropics (Suntharalingam et al., 2008; Berry et al., 2013) and also requires more uptake at high latitudes, especially in the NH.

To investigate the variation in the optimized fluxes of inversion S_u , we decompose the flux components as described in Section 2.1.4. The monthly fluxes and derived long-term trend are shown in ~~Figure ??~~. ~~Note that only the Supplementary Figure S7. The global flux was subsequently split into 8 regions, and the regional COS S_u fluxes analyzed for these regions~~ are shown in (Figure 6). Region NH1 (North America plus part of Pacific and Atlantic Oceans, orange) shows a negative "unknown" category (panel e) ~~has been optimized, and that yearly repeating ocean fluxes are not included in the figure. While there is no clear discernible trend in the prior biosphere and biomass burning fluxes, the prior anthropogenic emissions show a small increasing trend in 2003–2007 for COS and in 2010–2012 for CS₂. The optimized field shows an irregular seasonal cycle with substantial inter-annual variability, but no clear long-term trend. This indicates that the flux, indicating that more~~ sinks are needed. This likely points to an underestimation of the biosphere uptake in the prior, since this region (that is well constrained by observations) depicts a clear seasonal cycle in the optimized ~~"missing sources unknown" are likely partially related to fluxes with strong seasonal cycles~~ flux. A larger sink is also needed in NH2 (Europe, green) and NH3 (Asia, red), but of smaller magnitude than NH1. Tropical regions TR0–TR3 have similar trend and seasonality, and generally show a positive flux signal, with little seasonal cycle. This could represent an oceanic signal (underestimated emissions of COS or COS precursors in the prior), a signal from biomass burning, or an overestimated biosphere sink. The ocean-dominated region SH (blue) has a near neutral flux, with a seasonal cycle that shows higher emissions in local fall and early winter. In the next section, we will ~~therefore~~ explore the optimization of the ocean and biosphere fluxes.

3.2 Objective inversions

In this section we will discuss the results of inversions S1, S2, and S3. The resulting global budgets are compared to literature values in Table 4. In addition, χ^2 metrics and biases of the various inversions are reported in Table 5 for the NOAA surface network, the HIPPO campaigns, and the NOAA airborne profiles. Note that we also report results for optimizations that assimilated the HIPPO observations next to the NOAA surface data. The period of the analysed inversions is 2008–2010. The prior and posterior emission errors and error reduction of the different inversion scenarios are listed and discussed in Supplement Table S1.

The three inversions are all able to close the ~~gap in the~~ global COS budget, however, with very different budget terms (Table 4). Inversions S1 and S3 close the ~~COS gap in the~~ budget by a drastic reduction of the biosphere uptake in the tropics and more biosphere uptake at high latitudes. When the biosphere is not optimized (S2), the inversion enhances the CS₂ tropical oceanic source and reduces direct COS emissions from the high latitude oceans (Table 4). Both patterns lead to ~~enhanced release from the tropics reduced tropical biospheric uptake~~ and more uptake at high latitudes, as was found for inversion S_u .

Concerning the posterior fit to observations, none of the S1–S3 inversions performed like inversion S_u . The statistics in Table 5 show that S_u leads to the best fit to the assimilated observations, and only a small remaining bias. Inversions S1 and S3 show better χ^2 statistics and smaller biases than inversion S2, because it is difficult to fit continental NOAA stations (LEF, HFM, NWR, THD) only by optimizing ocean fluxes. However, S1 and S3 show a tendency to turn the tropical biosphere sink into a source, as shown in Figure 7, which depicts the posterior biosphere flux and flux increment for inversion S1. Note that while the uptake over high NH latitudes is enhanced, fluxes over regions in South America and over Indonesia have turned

into a source. This behaviour can be explained by the under-determined nature of the inverse problem: there are simply not enough observations [in the tropics](#) to constrain the tropical fluxes. Fast mixing in the tropics further complicates the detection of signals from the tropical biosphere using the NOAA surface network. Without additional observations it is therefore hard to unequivocally close the [gap in the](#) tropical COS budget. Currently, inversion S1 mostly assigns the missing sources to reduced
 375 biosphere uptake in the tropics, but the superior S_u inversion assigns the missing COS sources to a broad band in the tropics, without strong preference for land or ocean. Note that the behaviour of inversions S1, S2, and S3 is strongly driven by the predefined spatio-temporal patterns in the prior flux fields. [In section 3.4, we will revisit this issue.](#)

Although we currently cannot close the [gap in the](#) global COS budget with one specific known flux, it is instructive to explore the information content of [independent a separate set of](#) COS observations. In the next section, we will therefore evaluate the
 380 results of our inversions with HIPPO and NOAA airborne observations (Figure 1).

3.3 Evaluation with HIPPO and NOAA airborne profiles

From Table 5 it is clear that for all inversions the comparison to HIPPO observations is not very favourable. Most notably, the simulations with optimized fluxes show strong negative biases, and poor χ^2 statistics. However, Figure 8 shows that the inversions S1 and S_u (blue lines) largely improve the correspondence to HIPPO campaign 1 observations (red), relative to the
 385 prior simulation (black). The posterior simulations do capture the HIPPO observations much better. The remaining differences in the middle panels of Figure 8 show the general underestimation of the model. However, inversion S1 overestimates HIPPO in the southern tropics, likely caused by too large flux adjustments over South America, the region sampled by HIPPO campaign 1.

Interestingly, when the HIPPO observations are additionally assimilated in the inversion, biases are largely removed (Fig-
 390 ure 8, lower panels) while the correspondence to the NOAA surface network deteriorates only slightly (Table 5). Posterior χ^2 values for the HIPPO campaigns remain relatively poor, however, signalling too strict error settings or processes that are not properly modeled.

From the comparison with HIPPO we find that our state vector has enough flexibility to fit additional observations, and that the inversions are strongly observation-limited. Moreover, we find that the inversions based on only observations from
 395 the NOAA surface network tend to underestimate COS in the free troposphere. This is corroborated by observations from the NOAA airborne profiles, which are mostly collected over the US (see Figure 1). Figure 9 shows a comparison between profiles using results of inversion S1. Although most posterior profiles (blue) improve considerably compared to the prior simulation (black), they still underestimate observations (red) in the free troposphere. Note that the simulations based on inversion S1 correctly predict the draw-down of COS towards the surface for most measured profiles, and especially the match with the
 400 LEF site is very good at the surface, which confirms the performance of the inversion. If HIPPO observations are additionally assimilated (green), the agreement in the free troposphere slightly improves. For S1, χ^2 for the profile comparison reduces from 27.7 to 20.1, and the bias reduces from -13.9 to -9.7 pmol mol⁻¹ (Table 5). This confirms the low bias of the free troposphere COS mole fractions in simulations with fluxes that are optimized using both NOAA surface and HIPPO observations.

It is now clear that inversions using surface data from the available NOAA network sites will not be able to separate various source categories, and specifically not in the data-void tropics. In the next section we will therefore investigate the prospects of using satellite data to constrain fluxes.

3.4 Satellite validation

In Figure 10 we present a comparison between MIPAS, ACE-FTS, and co-sampled TM5 COS profiles. The latitude–height distributions of MIPAS, TM5 (convolved with the MIPAS AK) and ACE-FTS are shown in Figure 10(a-c). In Figure 10(d) we show averaged ACE-FTS, MIPAS and TM5 profiles, the latter two resulting from collocations with respect to ACE-FTS. The TM5 profiles shown are from the prior simulation (black), from inversion S1 (blue), and from inversion S1 with additional assimilation of HIPPO profiles (green), and averaged TM5 profiles. They are all convolved with the MIPAS AK of the prior (black) collocated with respect to ACE-FTS profiles.

In general, TM5 reproduces the observed pattern of COS well, but with lower values in the tropical up-welling region around 25 km altitude. The comparison between ACE-FTS and MIPAS is consistent with findings of Glatthor et al. (2017), who found that ACE-FTS is systematically lower in the UTLS region. Moreover, they found that MIPAS data showed no bias compared to MkIV and SPIRALE COS balloon profiles, which also exhibit higher COS values than ACE-FTS (Krysztofiak et al., 2015; Velazco et al., 2011). TM5 profiles, after convolution with the MIPAS AK, are in between MIPAS and ACE-FTS. Prior TM5 profiles (black) show highest values around the tropopause. Again, TM5 profiles optimized by HIPPO and NOAA observation (green dashed line in Figure 10d) show a slight increase in the upper troposphere compared to the optimization with only NOAA surface-site data (blue dashed line).

To compare the different inversions with respect to the simulated latitude–longitude distribution, Figure 11 shows a comparison of COS between TM5 results–are inversions and MIPAS on 250 hPa in June to August. Similar results on 250 hPa from September to November and on 150 hPa from June to August are shown in supplement (Figure S8 and Figure S9). MIPAS COS represents a 2002–2011 average taken from Glatthor et al. (2017). TM5 results have been averaged over 2008–2010. The distributions of COS in all inversions match relatively well with MIPAS. Note, however, that we adjusted the TM5 results by +25 pmol mol^{−1} to match the colorscale of MIPAS. The COS distribution from the prior simulation correctly simulates low COS over the Amazon and Africa, but is clearly too high over Northern latitudes. This latter aspect is partly solved by the inversions. If we concentrate on the observed COS minimum over the Atlantic, Africa and the Amazon, inversions S1 and S3 shift this minimum to the east, consistent with the COS biosphere flux increment shown in Figure 7 for S1. Inversions S₀ and S2 exhibit a better comparison with MIPAS, suggesting that the large increments of the tropical biosphere over South America (Figure 7) are unrealistic. However, assigning the missing tropical source totally to ocean emissions (S2) appears to overestimate the COS drawdown over the Amazon.

TM5 results are also compared to the nadir viewing TES instrument. To this end, COS columns of TM5 (convolved with the TES AK, see Equation 1) and TES are averaged in 20 latitudinal bins between 32°S and 32°N. Outside this latitude band, TES observations become too noisy for a reasonable comparison. Comparisons are shown for the months March, June, September and December in Figure 12, based on inversion S1, and averaged over the years 2008–2011. This comparison

shows that the prior simulation is too high in the tropical latitudes and on the NH (e.g., June, September, and December). After assimilation, the agreement with TES improves, but now a general underestimate can be observed. The inversion in which also the HIPPO observations are assimilated bring the simulated mole fractions closer to TES (except for September), confirming our earlier findings based on the airborne observation. Thus, although the TES-derived columns are rather noisy, they offer good perspective to better constrain the COS budget in the tropics. Due to the sensitivity of TES to COS in the middle troposphere (Kuai et al., 2015), the assimilation of TES in our 4DVAR system might be able to differentiate between the biosphere and ocean signal, something that turned out to be difficult using NOAA surface observations only.

3.5 Discussion

In this study we have presented inversions focused on the closure of the global COS budget. In general, our inversion modelling framework based on the TM5-4DVAR system is quite well capable to close the gap in the global budget (e.g., inversion S_u , S1 and S3) and to optimize flux fields such that surface observations are well reproduced. However, due to the lack of observations, we are unable to unambiguously assign the missing COS sources to either missing ocean emissions or to reduced tropical uptake by the biosphere. Firstly, the total number of observations remains relatively small, which leads to an under-determined inversion problem. Secondly, there are no observational sites that sample air masses from tropical Africa, South America, and South East Asia, which are regions with important COS fluxes. An important next step will therefore be the utilization of satellite data in future inverse modelling studies. In the current study, we did not include all exchange fluxes that are reported in literature (Whelan et al., 2018). In general, we find that our inversions still underestimate COS in the free troposphere. Here, there might be a role for volcanic emissions ($25\text{--}42\text{ Gg S a}^{-1}$ (Whelan et al., 2018)), or 'unnoticed' tropical sources like wetland exchange ($\sim 150\text{ to }290\text{ Gg S a}^{-1}$ (Whelan et al., 2018)). Volcanic emissions are important to mitigate the stratospheric aerosol loading in the stratosphere (Sheng et al., 2015) and might be able to reduce the gap between modelled COS by TM5 and measurements. Alternatively, missing COS could come from an atmospheric oxidation process that converts CS_2 or DMS to COS. We did not find strong evidence for enhanced CS_2 emissions from tropical oceans in our S1 inversion, although inversion S2 produced reasonable COS simulations by optimizing only COS and CS_2 emissions from the ocean. Moreover, our "best" S_u inversion produced a flux field that indicated enhanced tropical sources over both land and ocean (Figure 5). Thus, field studies that address tropical COS exchange processes are urgently needed (Lennartz et al., 2020).

We have also considered some variations in our modelling setup. A unique approach of our study is the inclusion of CS_2 and DMS as COS precursors. We tested the effect of emitting CS_2 ocean and anthropogenic sources directly as COS in an additional forward model simulation. As shown in Figure S10, COS mole fractions would become significantly larger close to CS_2 emission hot spots in Asia, Europe and the US. At selected stations (LEF in the US and MHD in Europe, Figure S10 a and b), we observe COS mole fractions that are up to 40 pmol mol^{-1} higher during events where emitted CS_2 is advected to the station. Some ambiguity has been introduced about the CS_2 lifetime (Khan et al., 2017). In our S_u inversion, the lifetime of CS_2 is estimated as 9.4 days (CS_2 burden divided by CS_2 loss by OH), substantially longer than the ≈ 3 day lifetime mentioned in Khan et al. (2017). Future work should be based on the rate recommendations in Sander et al. (2006). Thus, we conclude that inclusion of CS_2 as a separate tracer is important if we want to understand emissions of CS_2 and COS, which have

distinctly different spatial patterns (e.g. see Supplementary Figure S4). Regarding DMS as COS precursor, we have evaluated its importance by performing a NO-DMS inversion, in which DMS as a tracer was removed and the 162 Gg S a⁻¹ DMS source was added to the COS "unknown flux" in inversion S_u. In Supplement Figure S11, it can be seen that the NO-DMS inversion shows larger adjustment over both oceans and continents, but that the pattern remains comparable to inversion S_u.

The use of COS as a proxy for gross primary productivity on a global scale needs a better level of understanding of the biosphere flux. Here we used monthly prior flux fields calculated with the SiB4 model (Berry et al., 2013) in which soil exchange and vegetation uptake are combined. In future studies, we might need a better prior description of this important global COS sink. For instance, recent studies (Ogée et al., 2016; Sun et al., 2018; Meredith et al., 2019; Spielmann et al., 2020) stress the importance of the soil-atmosphere COS exchange. Our inversions S1 and S3 calculate large increments in the biosphere exchange (Figure 7), with general less uptake in the tropics (turning the flux even into a COS source) and enhanced uptake in the NH high latitudes. Quantitatively, the COS uptake is reduced from a prior value of 1053 Gg S a⁻¹ to 557 Gg S a⁻¹ to close the gap in the COS budget. While we seriously question the validity of this result given the fact that most flux adjustments are projected in the data-void tropics, it is still instructive to consider the feedback of the atmospheric COS mole fractions on COS uptake. Since biosphere models operate mostly uncoupled to atmospheric transport models, we used a fixed mole fraction of 500 pmol mol⁻¹ to construct the prior biosphere fluxes. However, observations clearly show a large drawdown of COS near the surface (Hilton et al., 2017; Spielmann et al., 2020) (Campbell et al., 2008; Hilton et al., 2017; Spielmann et al., 2020; Berkelhammer et al., 2020). We therefore explored the calculations in SiB4 and found that biosphere flux should scale linearly with atmospheric COS mole fractions (Berry et al., 2013). To estimate the potential impact of reduced mole fractions at the surface on the biosphere flux, we corrected the monthly SiB4 fluxes as:

$$f_{\text{biOSP,cor}} = f_{\text{biOSP}} \frac{y(\text{COS})}{500 \text{ pmol mol}^{-1}}, \quad (11)$$

where f_{biOSP} and $f_{\text{biOSP,cor}}$ are the original and corrected monthly biosphere fluxes on the TM5 grid, and $y(\text{COS})$ is the monthly mean COS mole fractions (pmol mol⁻¹) in the first model layer (approximately 50 m) from inversion S_u. This simple correction, based on monthly mean fields, changes the biosphere sink from ~~-1053~~-1053 Gg S a⁻¹ to ~~-851~~-851 Gg S a⁻¹, an update of 202 Gg S a⁻¹ (Supplementary Figures S12, S13 and S14), and closer to the 738 Gg S a⁻¹ reported by Berry et al. (2013). Interestingly, the corrected flux is strongly reduced over regions with an active tropical biosphere, in line with results from inversions S1 and S3. This indicates that uptake of COS should be treated as a first order loss process, and that the SiB4 prior fields based on fixed atmospheric mole fractions of 500 pmol mol⁻¹ likely overestimate COS uptake. However, such an approach makes the optimization problem non-linear. This, and the challenge of assimilating satellite observations, will be the subject of future studies.

4 Conclusions

In this study, we have implemented an inverse modelling framework for COS, coupled to the budgets of CS₂ and DMS. Inversions using the NOAA surface observation network have been evaluated with observations from HIPPO, airborne observations, and satellite products. Conclusions are:

- In line with earlier studies, our inversions point to missing sources in the tropics and missing sinks at high latitudes. ~~We identify that the flux field that closes the budget exhibits an irregular seasonal cycle~~ With seasonal decomposition of the optimized unknown COS flux, it is found that the missing sources show regional seasonality, indicating regional source or sink impacts. Whether the missing sources in the tropics originate from the land or ocean cannot be determined currently ~~because of a lack of~~ because of a lack of observations in the tropics ~~are sparse. Part of the tropical sources can be explained by the dependence of COS uptake on atmospheric mole fractions.~~
- Simulations that are optimized by only NOAA surface observations from 14 sites lack information about COS in the free troposphere. When the short-term HIPPO aircraft sampling program is used as an additional data source in the inversions, the comparison to NOAA airborne observations and satellite products generally improves.
- Comparison between TM5 inversions and satellite data shows that COS in the model is systematically lower than MIPAS or TES, and inversions reproduced tropospheric COS spatial distribution well, specifically for inversions S_u and S2. These comparisons indicate that the missing tropical source likely originates from a combination of underestimated ocean emissions, and overestimated biosphere uptake. Part of the tropical sources can be explained by the dependence of COS uptake on atmospheric mole fractions.
- Future improvements are expected from the assimilation of satellite data and better prior descriptions of the ocean and biosphere fluxes.

Our future plan is therefore to assimilate satellite data into our 4DVAR inverse modelling system to have better constraints on COS in the free troposphere and lower stratosphere. Other developments target the coupling of COS and CO₂ in a shared inverse modelling system, with the aim to better constrain gross primary productivity.

Code availability. Codes are available on request

Author contributions. JM and MK designed the study. JM implemented the method and carried out numerical experiments. LMJK and AC performed SiB4 model experiments. SM and EA provided observations from the NOAA surface network, HIPPO and airborne observations. JW and LK provided the TES–Aura satellite product and helped with the technical discussion on the use of satellite products. NG provided the MIPAS satellite data product with averaging kernel and helped with the discussion. JM and MK wrote the manuscript with contributions from all co-authors.

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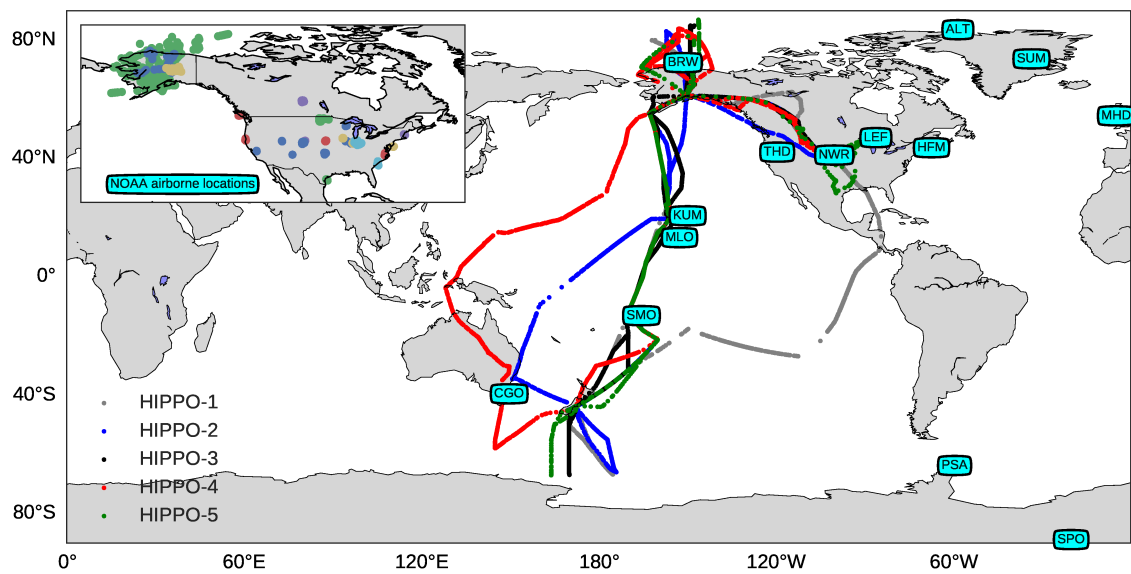


Figure 1. Geographical locations of the NOAA ground-based observations (shown in boxes), the five HIPPO campaign tracks, and the NOAA profile program (inset). Note that there are no NOAA surface stations located in Asia, South America, ~~Africa~~ or ~~Europe~~ Africa.

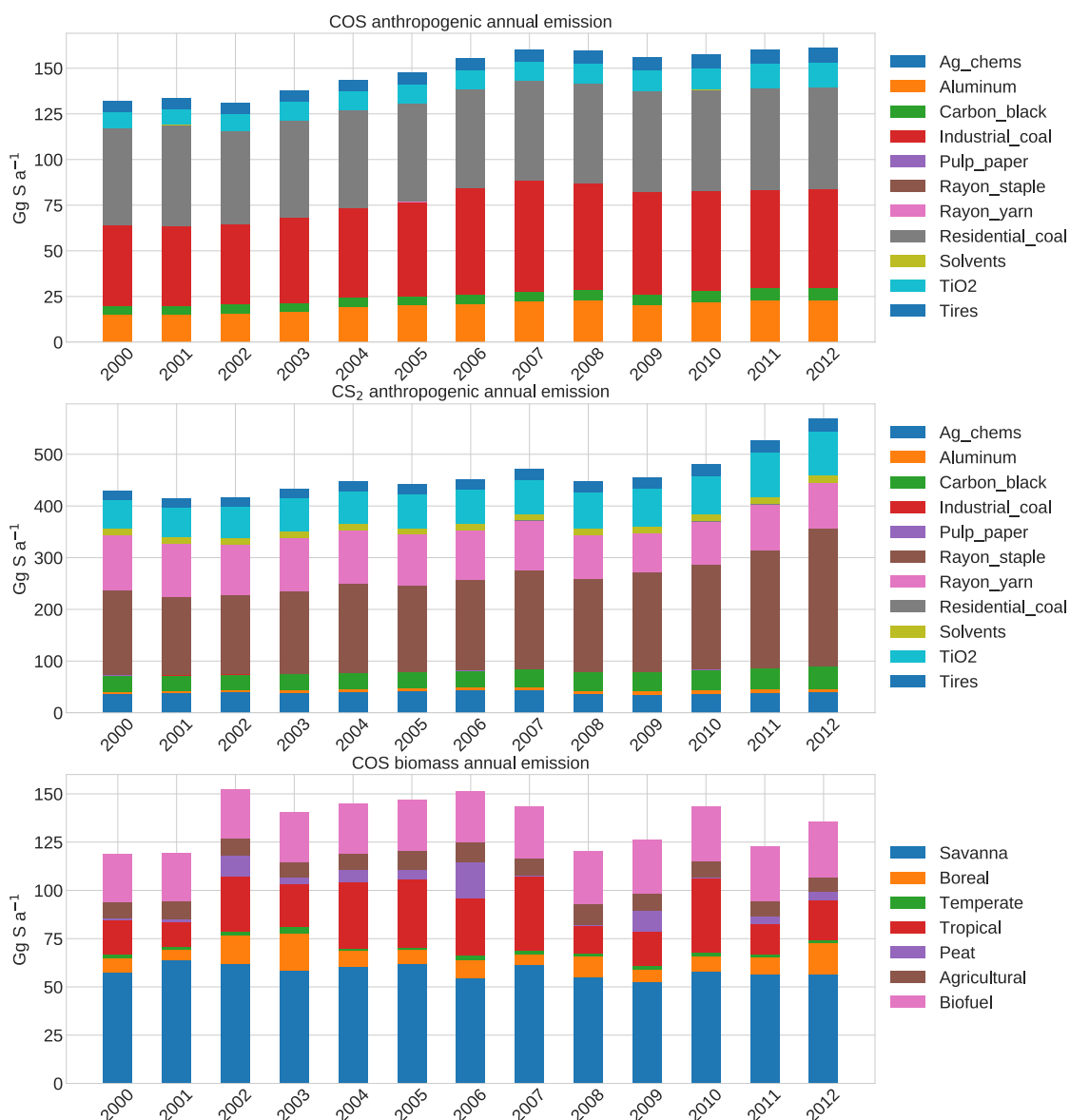


Figure 2. Yearly anthropogenic emissions of COS and CS₂ and COS biomass burning emissions in the period 2000 to 2012. We disentangled the emissions reported in Zumkehr et al. (2018) into COS and CS₂ emissions using their reported yield of 0.87 (see main text). Biomass burning emissions are calculated based on the GFED 4.1 biomass burning inventory and the CEDS biofuel emission inventory (see main text).

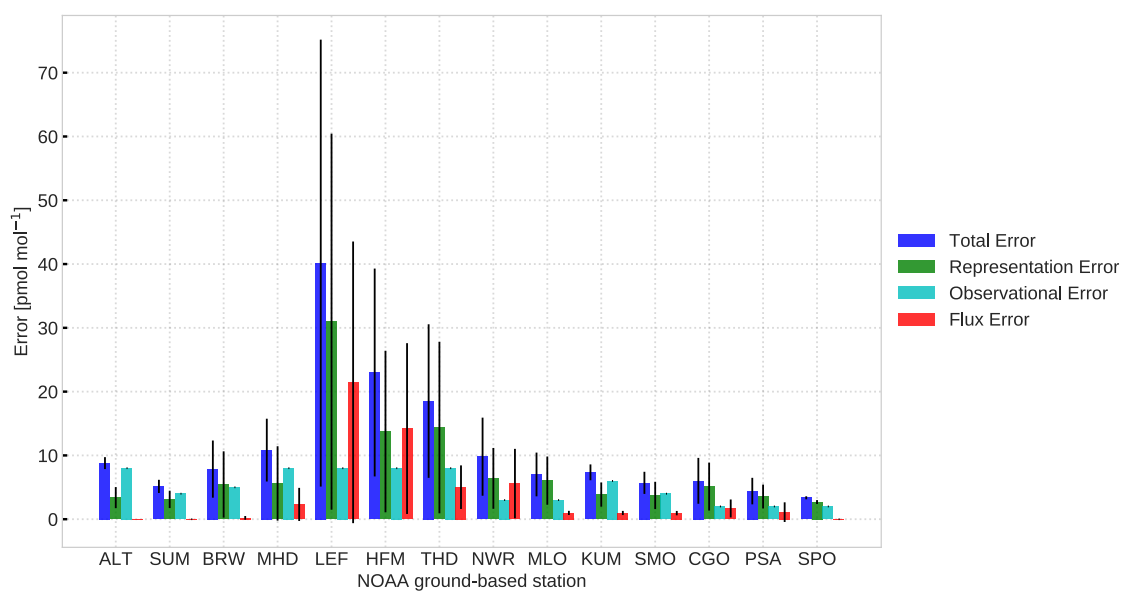


Figure 3. Error analysis for NOAA stations. Black error bars represent the time-variations of the errors over a 3-year period (2008–2010). For ALT, SPO and SUM, the flux-related errors are close to zero and not shown. Stations are ordered from the North Pole to the South Pole.

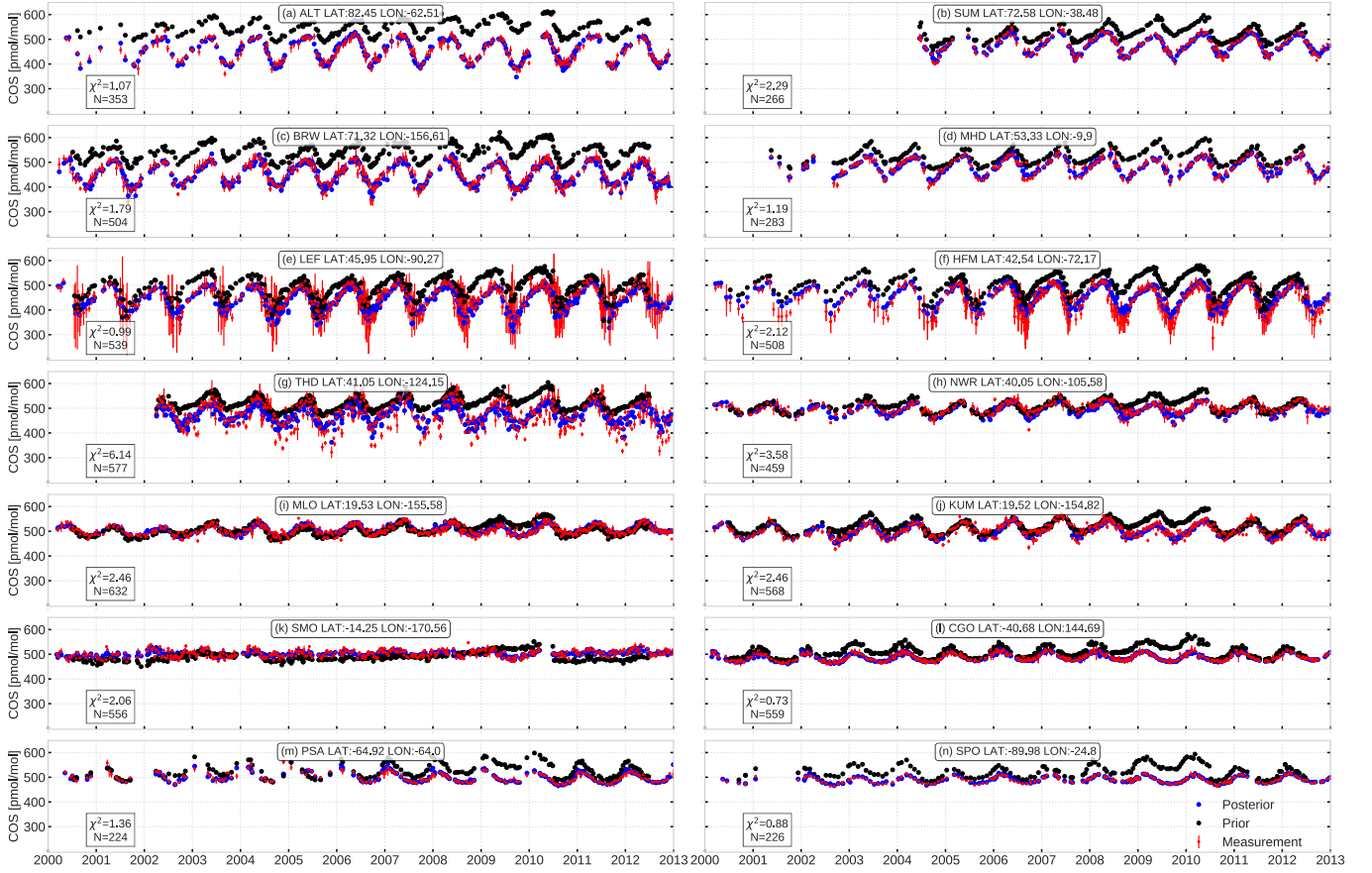


Figure 4. COS prior and posterior comparison at NOAA stations. Red dots and bars are NOAA measurements with errors. Blue and black dots represent the posterior and prior simulation, respectively. Results are shown for inversion S_u in which only the "unknown" emission category is optimized.

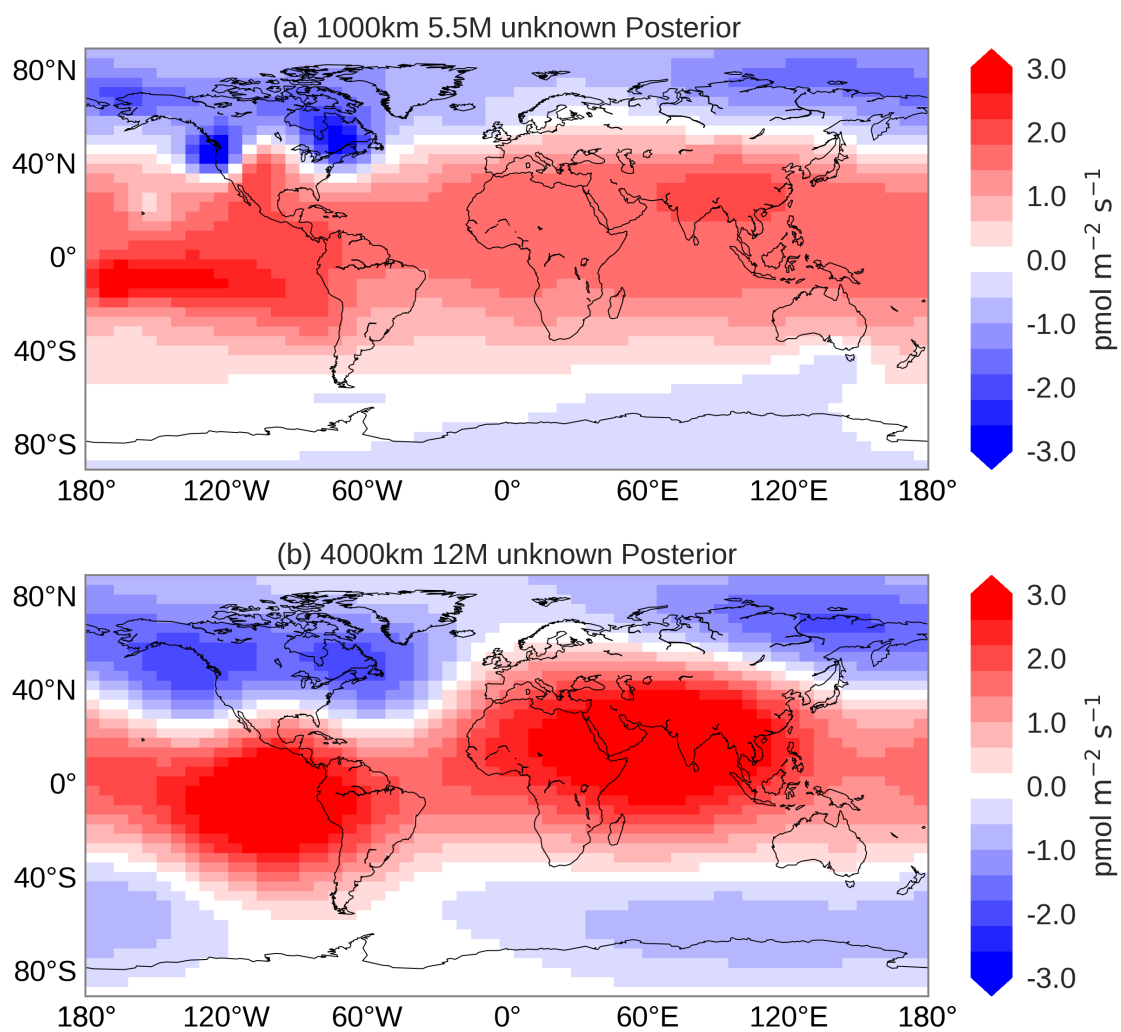


Figure 5. Optimized emission pattern of the "unknown" field of inversion S_u for different settings of the spatio-temporal correlation lengths (a) Spatial correlation of 1000 km and temporal correlation of 5.5 months (b) Spatial correlation of 4000 km and temporal correlation of 12 months. Results are averages over 2008–2011.

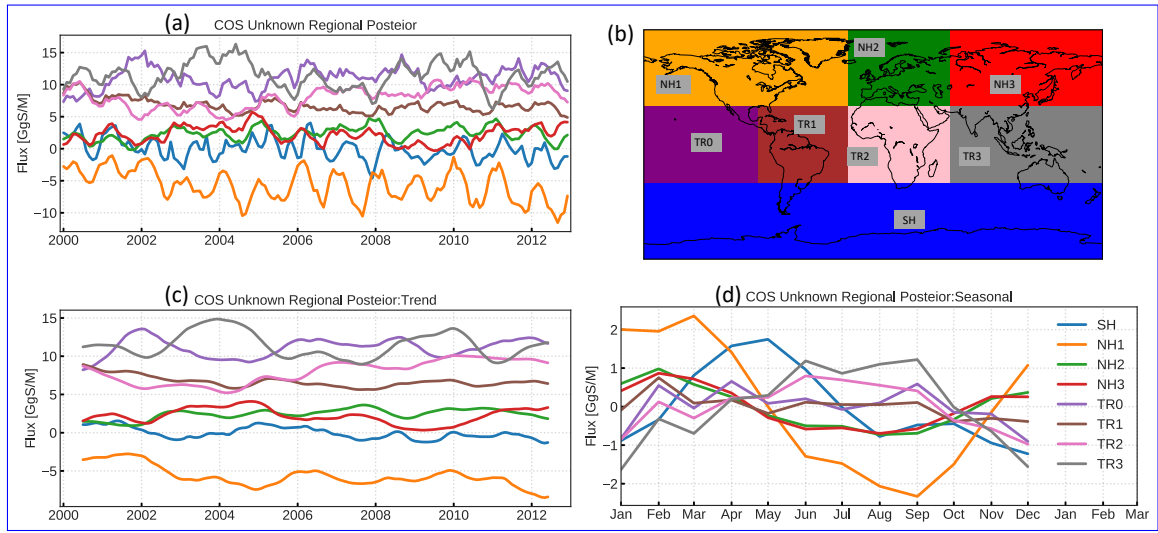


Figure 6. Monthly-exchange-Regional analysis of multi-year optimized COS fluxes related to of inversion S_u : (a) prior-biosphere-posterior flux per region (b) anthropogenic COS emissions regions over which the posterior flux is analysed (c) COS from anthropogenic CS_2 -trend in the decomposed signal (d) biomass-burning (e)-optimized-"unknown"-emissions seasonal signal in the decomposed signal. Monthly-global emissions Note that region colors in (fluxesb) are shown-used in gray-panels (a), (c) and the-trend-is-shown-in-blue(d).

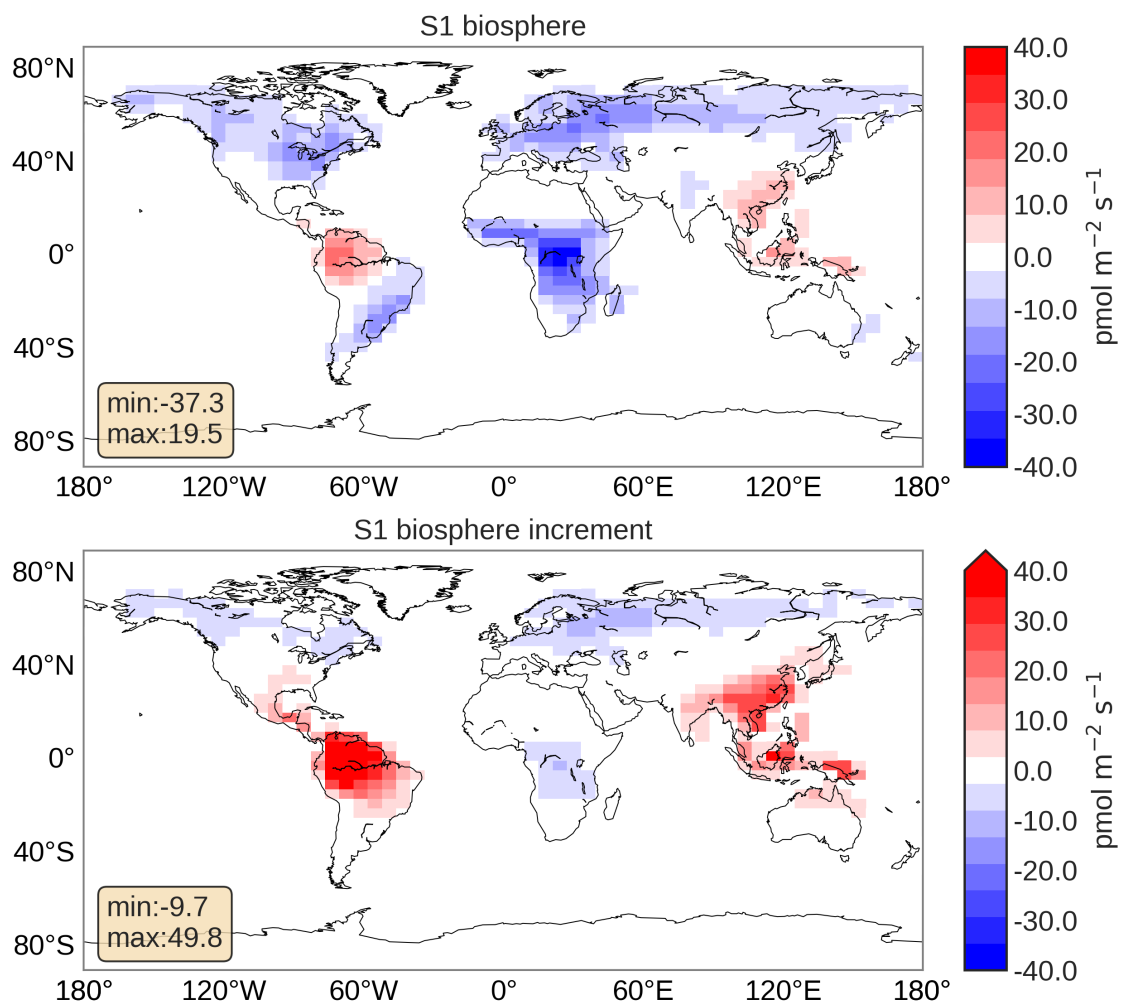


Figure 7. Posterior biosphere flux from inversion S1 and increment (posterior-prior). The fluxes represent 3-year (2008–2010) averages with removal of 6 month spin-up and spin-down periods. The maximum and minimum flux values are given in the boxes.

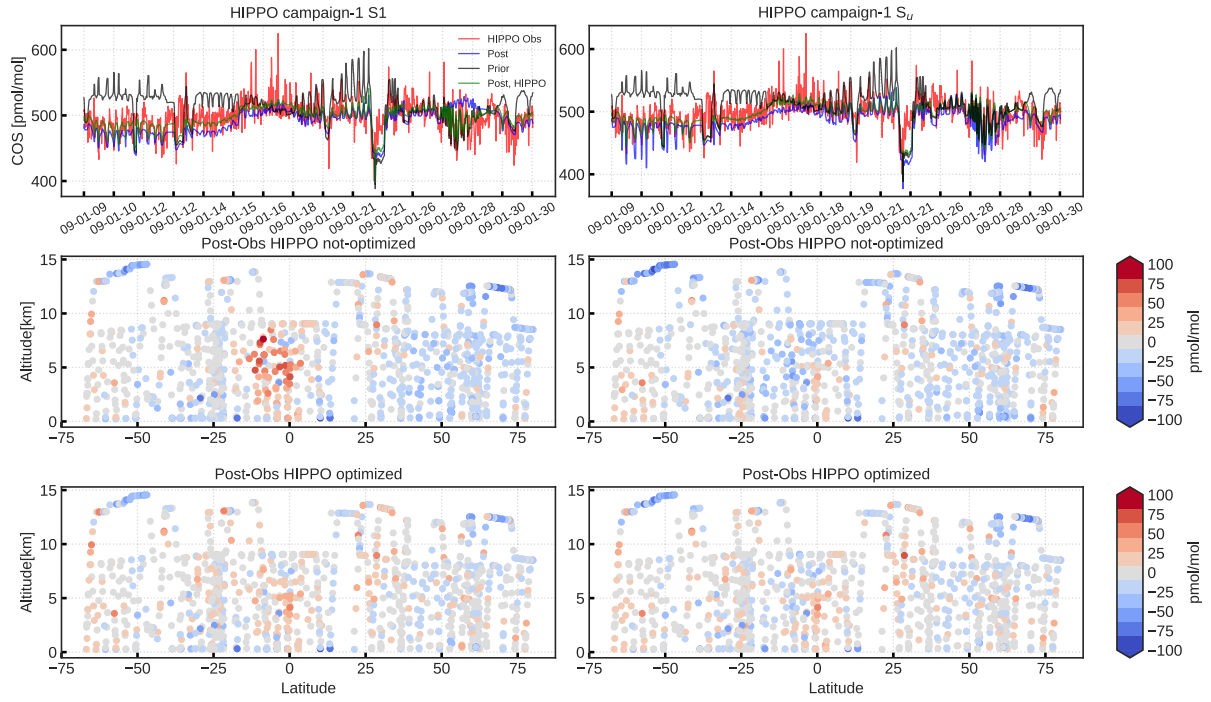


Figure 8. HIPPO campaign 1 COS observations compared to results from inversions S1 (left) and S_u (right). The first row shows time series of HIPPO observations (red), prior (black), posterior (blue), and posterior with HIPPO observations assimilated (green). The middle and bottom rows show model minus observations in a latitude-height plot for inversions with and without assimilating HIPPO observations, respectively.

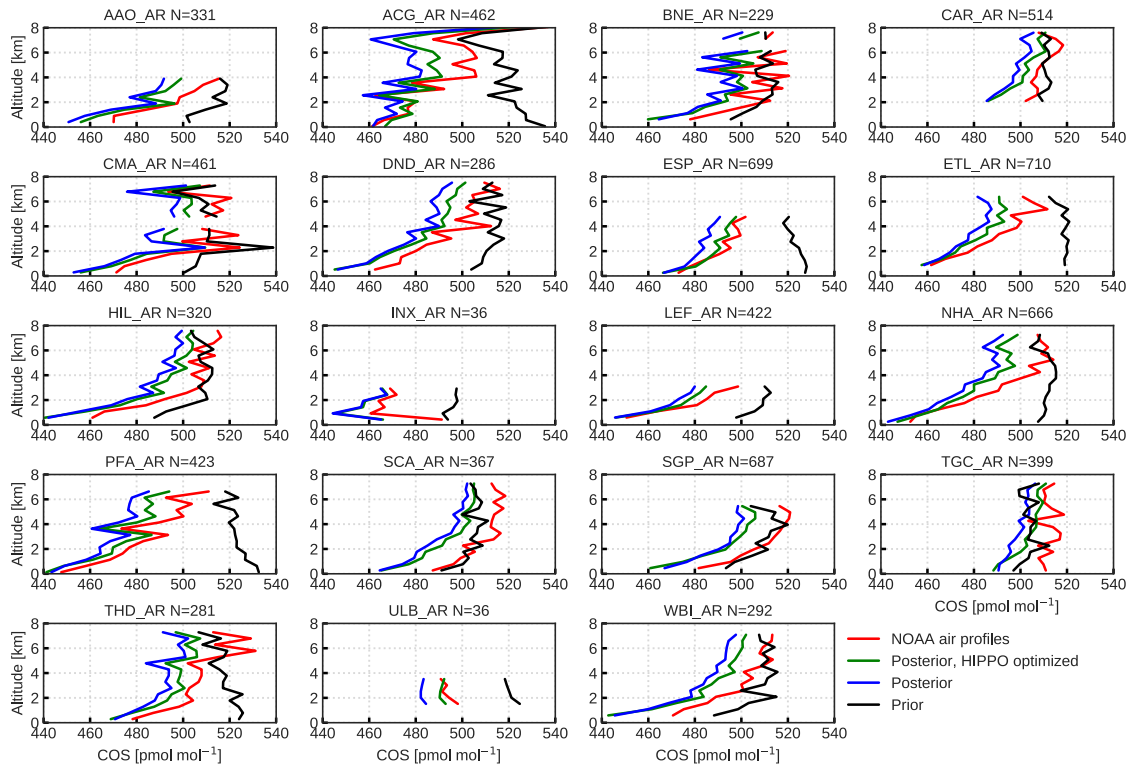


Figure 9. Prior (black) and posterior (blue) profiles of inversion S1, compared to NOAA aircraft profiles (red). Location (see Figure 1) and number of observations are mentioned in the caption. The green lines are results from an inversion in which, next to NOAA surface observations, also HIPPO observations are assimilated. Note that the profiles are from all seasons.

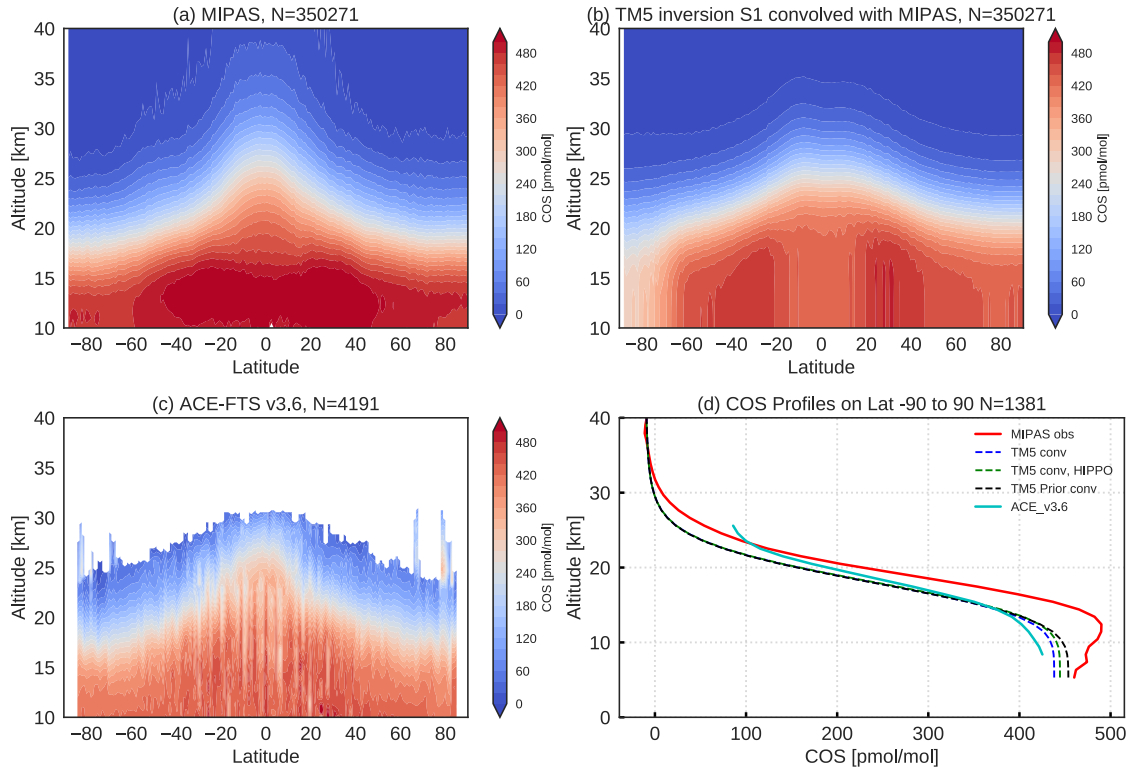


Figure 10. Comparison of MIPAS and ACE-FTS v3.6 with TM5 results from ~~inversion S1 in inversions for~~ 2009. (a) latitude–height contour plot of MIPAS; (b) TM5 S1 convolved with the MIPAS AK; (c) ~~ACE-FTS~~ ACE-FTS profiles; (d) average of collocated profiles for MIPAS (red), TM5 convolved with MIPAS AK from inversion S1 (blue), TM5 convolved from inversion S1 (with HIPPO observations assimilated) (green), TM5 convolved prior (black), and ACE-FTS. In (d) TM5 and MIPAS profiles are collocated with respect to ACE-FTS profiles within a temporal offset of 6 hr and a spatial distance within 5 degrees. The number of collocated profiles is 1381.

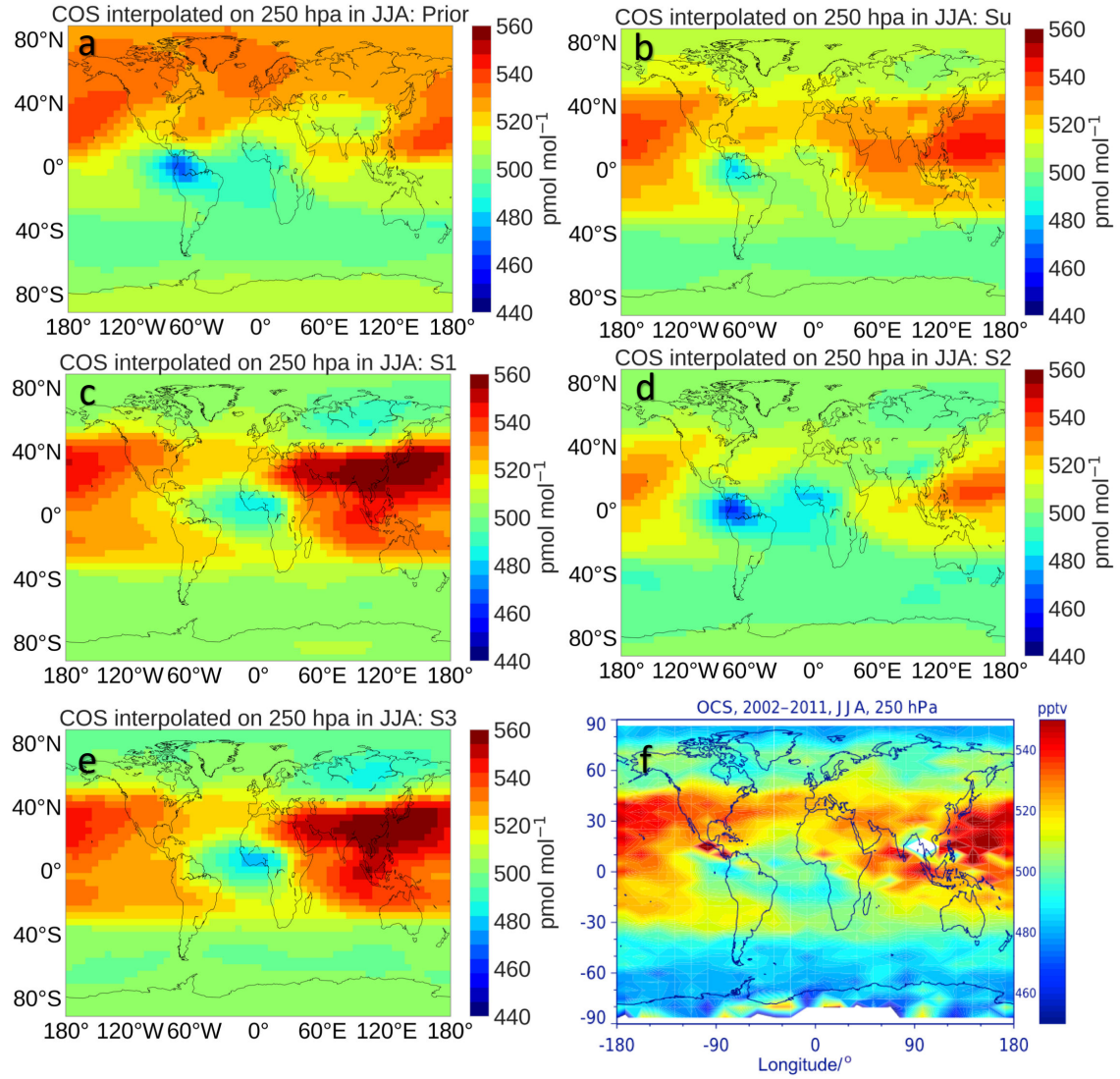


Figure 11. COS mole fraction comparison of MIPAS and TM5 inversions on 250 hPa in June to August. (a–e) are TM5 prior, and inversions S_u , $S1$, $S2$, $S3$, respectively. (f) is captured from figure 11 in (Glatthor et al., 2017). TM5 results represent a 2008–2010 average, and MIPAS is averaged over 2002–2011. Because TM5 results are systematically lower than MIPAS, 25 pmol mol^{-1} is added to the TM5 results for a better visual comparison.

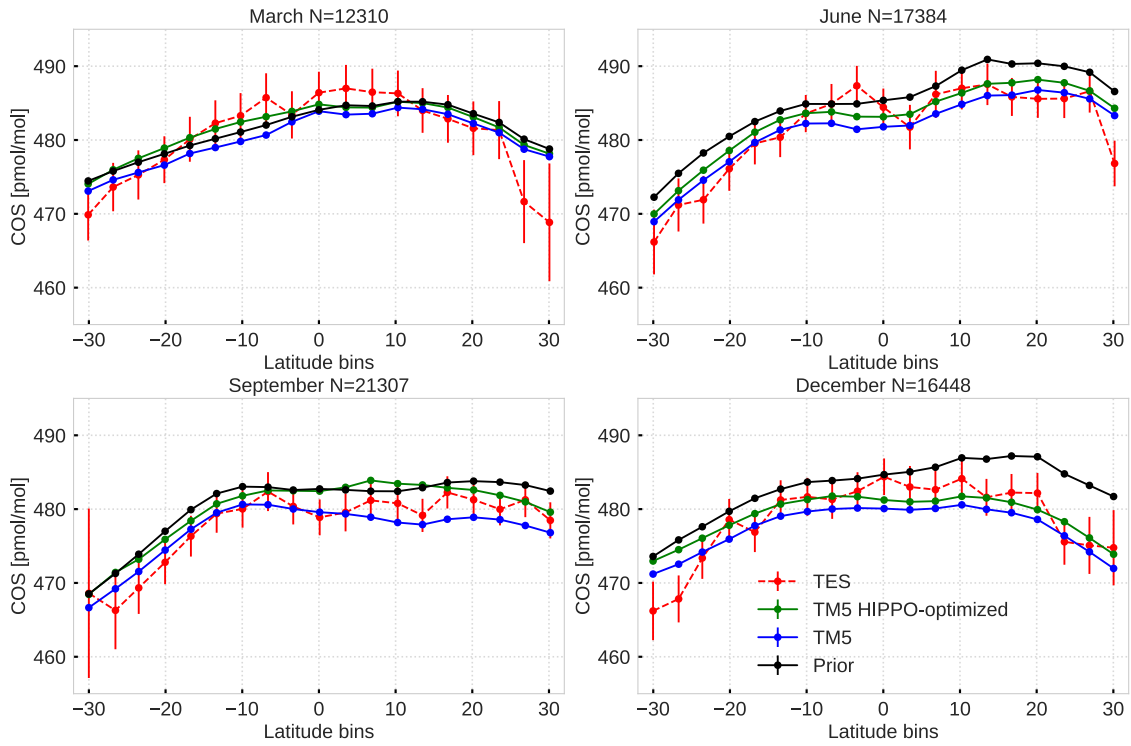


Figure 12. Column averaged COS mole fractions sampled by TES (red), model prior (black), model posterior (blue) and posterior with HIPPO observations assimilated (green) for March, June, September and December. The columns are averaged over 2008–2010 in 20 latitudinal bins from 32°S to 32°N. The result of inversion S1 is shown. Error bars on TES represent variability in the measurements and the number of observations is given in the caption. Variability is mostly determined by measurement noise.

Table 1. The split of anthropogenic emissions in the different categories and between COS and CS₂ based on Zumkehr et al. (2018). Note that we used a CS₂ to COS molar yield of 0.87 and that CS₂ contains two S atoms. Averages over 2000–2012 are presented.

Emission Type	Total COS	Fraction COS ^a	Direct COS	Direct CS ₂
	Gg S a ⁻¹	%	Gg S a ⁻¹	Gg S a ⁻¹
Agricultural Chemicals	16.9	0.0	0.0	38.9
Aluminum Smelting	22.2	88.2	19.6	6.0
Industrial Coal	52.1	99.5	51.8	0.7
Residential Coal	54.0	100.0	54.0	0.0
Industrial Solvents	5.4	0.0	0.0	12.5
Carbon Black	19.7	26.5	5.2	33.3
Titanium Dioxide	39.4	26.5	10.5	66.6
Pulp & Paper	0.1	3.2	0.0	0.3
Rayon Yarn	41.1	0.0	0.0	94.6
Rayon Staple	77.3	0.0	0.0	177.7
Tires	15.1	43.0	6.5	19.8
Total Anthropogenic	343.3	-	147.5	450.2

^a The fraction of COS is calculated based on the COS to CS₂ emission ratio reported in Table 1 of Lee and Brimblecombe (2016).

Table 2. Biomass burning emission factors used in converting COS emissions. EF COS denotes the COS emission factor from dry mass in units g kg^{-1} COS per dry mass, and EF CO denotes the CO emission factor in g kg^{-1} CO per dry mass. Emission factors were taken from Andreae (2019).

	EF COS g kg^{-1} COS per dry mass	EF CO g kg^{-1} CO per dry mass
Savanna and grassland	0.038	-
Tropical forest	0.078	-
Temperate forest	0.035	-
Boreal forest	0.058	-
Peat fires	0.110	-
Agricultural waste burning	0.059	-
Biofuel burning without dung	0.017	83
Biofuel burning with dung	0.210	89

Table 3. Names and error settings of the inversions performed in this study. The values correspond to grid-scale errors. Monthly flux fields are optimized using spatial and temporal correlation lengths of 4000 km and 12 months, except for inversion S_u , in which multiple settings are explored.

	Biosphere	Ocean COS	Ocean CS ₂	Biomass burning	Anthropogenic COS and CS ₂	"Unknown"
S_u	-	-	-	-	-	100%
S1	50%	50%	50%	10%	10%	-
S2	-	50%	50%	-	-	-
S3	50%	-	-	-	-	-

Table 4. Results from inversions S_u , S1, S2, and S3 compared to published global COS budgets

COS Budget (Gg S a^{-1})	Kettle2002 ^a	Montzka2007 ^b	Berry2013 ^c	Kuai2015 ^d	Our prior	S_u	S1	S2	S3
Direct oceanic COS	41	40	39	41	40	40	-18	22	40
Indirect oceanic CS_2 as COS	84	240	81	83	81	81	96	499	81
Indirect oceanic DMS as COS	154		156	155	156	156	156	156	156
Direct anthropogenic COS	64	64	64	62	155	155	153	156	155
Indirect anthropogenic CS_2 as COS	116	-	116	113	188	188	188	189	188
Indirect anthropogenic DMS as COS	1	-	1	0	6	6	6	6	6
Biomass burning	38	106	136	49	136	136	124	136	128
Additional ocean flux	-	-	600	559	-	-	-	-	-
Anoxic soils	26	66	-	-	-	-	-	-	-
Sources	523	516	1193	1062	762	762	705	1163	754
Destruction by OH	-94	-96	-101	-111	-101	-101	-103	-101	-101
Destruction by O	-11	-11	-	-	-	-	-	-	-
Destruction by photolysis	-16	-16	-	-	-40	-40	-40	-40	-40
Uptake by plants	-238	-1115	-738	-775	-1053	-1053	-557	-1053	-613
Uptake by soil	-130	-127	-355	-176					
Sinks	-489	-1365	-1194	-1062	-1194	-1194	-700	-1194	-754
Unknown	-	-	-	-	432	425	-	-	-
Net total	34	-849	-2	0	0	-6	5	-31	0

^a Kettle et al. (2002)

^b Table 2 from Montzka et al. (2007)

^c Berry et al. (2013)

^d Kuai et al. (2015)

Table 5. χ^2 metrics and mean biases for the different inversion scenarios. Statistics are shown for the NOAA surface stations, the HIPPO campaigns, and the NOAA airborne profiles. Biases are given in pmol mol^{-1} .

Inversion scenario	HIPPO optimized ^a	Metric	HIPPO	NOAA surface	NOAA airborne
S_u	No	χ^2	40.7	1.9	26.0
	No	Bias	-13.9	0.0	-12.4
	Yes	χ^2	4.7	2.5	17.3
	Yes	Bias	-1.1	1.5	-8.3
S1	No	χ^2	43.8	2.4	27.7
	No	Bias	-12.0	-0.4	-13.8
	Yes	χ^2	4.8	2.9	20.1
	Yes	Bias	-1.3	1.3	-9.7
S2	No	χ^2	54.2	4.9	48.2
	No	Bias	-19.4	1.5	-16.7
	Yes	χ^2	6.3	5.9	27.0
	Yes	Bias	-4.6	7.5	-5.9
S3	No	χ^2	43.3	2.5	27.5
	No	Bias	-12.3	-0.2	-14.3
	Yes	χ^2	5.0	3.2	21.1
	Yes	Bias	-1.4	1.6	-10.5

^a If HIPPO is not optimized, only NOAA surface data is assimilated in inversions. If HIPPO is optimized, both NOAA surface data and HIPPO are assimilated in inversions. NOAA airborne data is only used for validation.

Supplement of

Inverse modelling of carbonyl sulfide: implementation, evaluation and implications for the global budget

Jin Ma et al.

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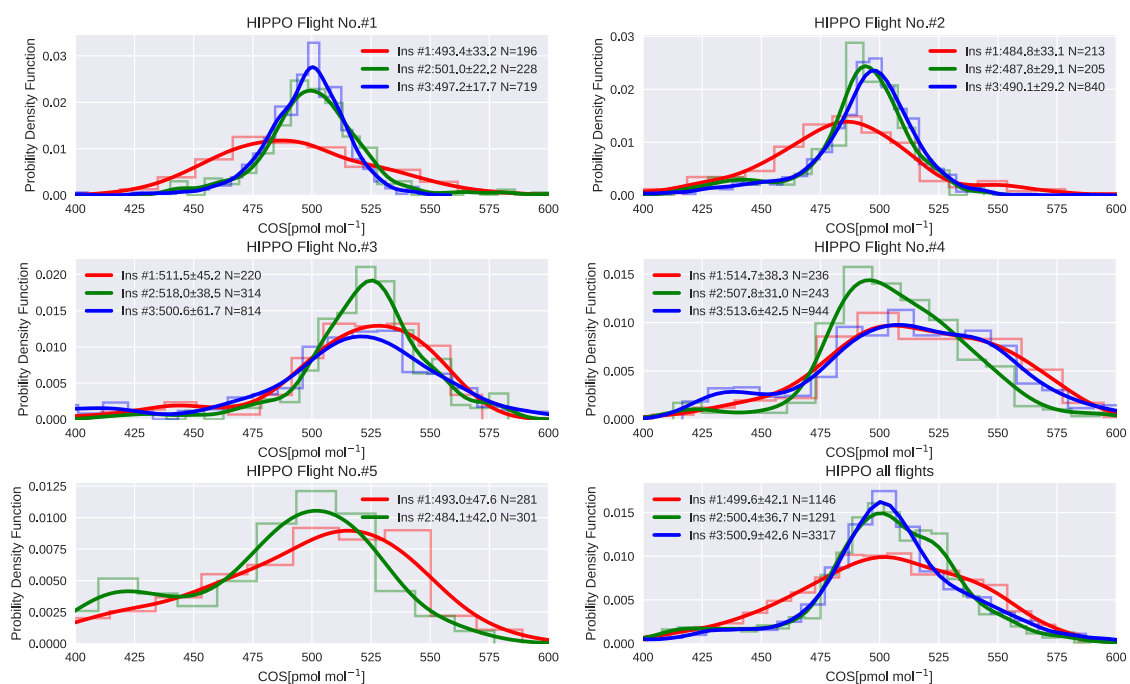


Figure S1. COS mixing ratio distribution measured by HIPPO instruments 1–3. The measurements of COS were taken by 3 different instruments. The last panel represent the averages of all HIPPO flights. The mean and number of measurements of each flight are shown in legends.

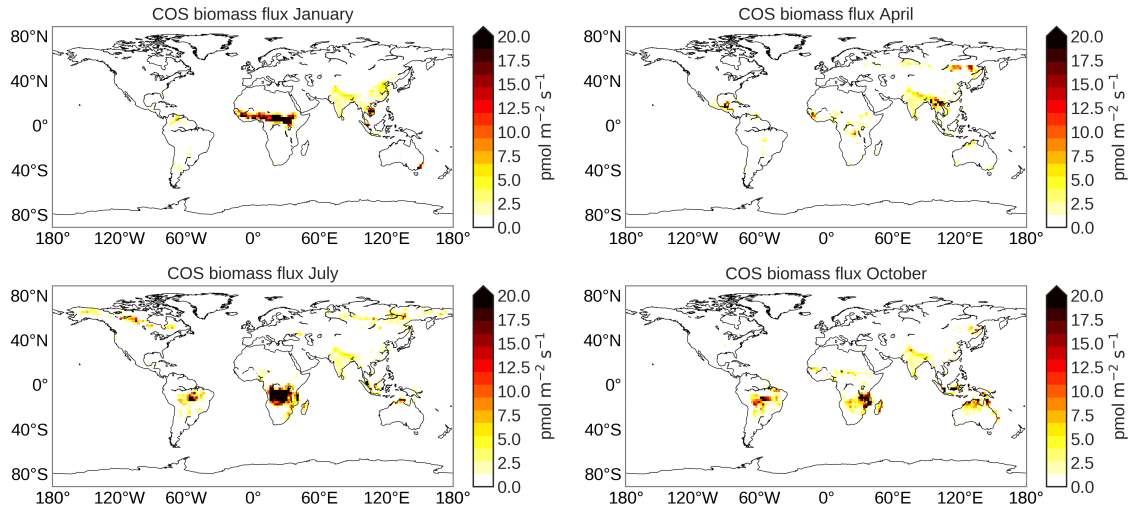


Figure S2. Spatial distribution of prior COS biomass burning fluxes for January, April, July, and October. Note that we use an extended colorbar due to hot spot emissions. The fluxes are averaged over the years 2000–2012.

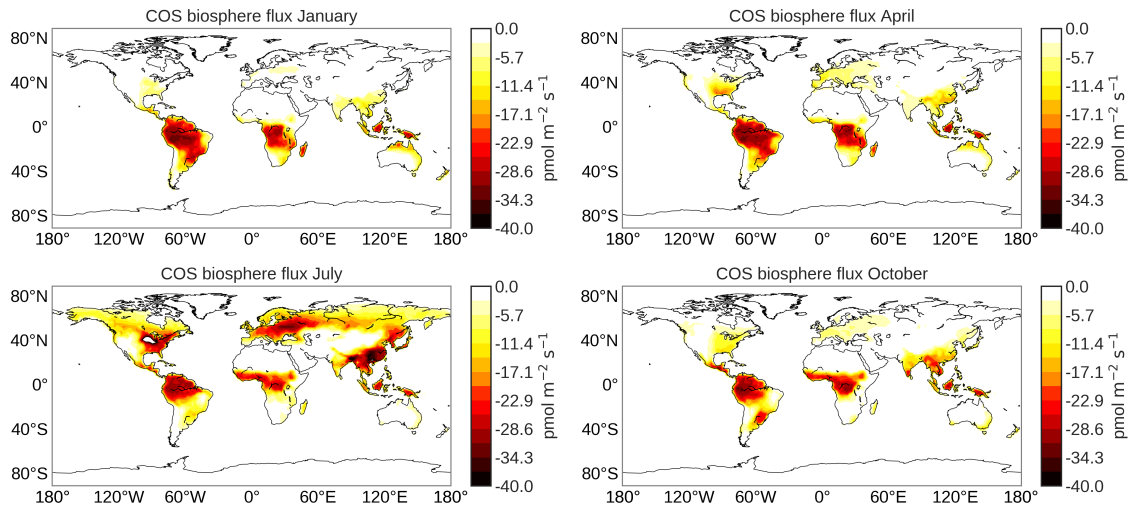


Figure S3. Spatial distribution of prior COS biosphere fluxes for January, April, July, and October. The fluxes are averaged over the years 2000–2012.

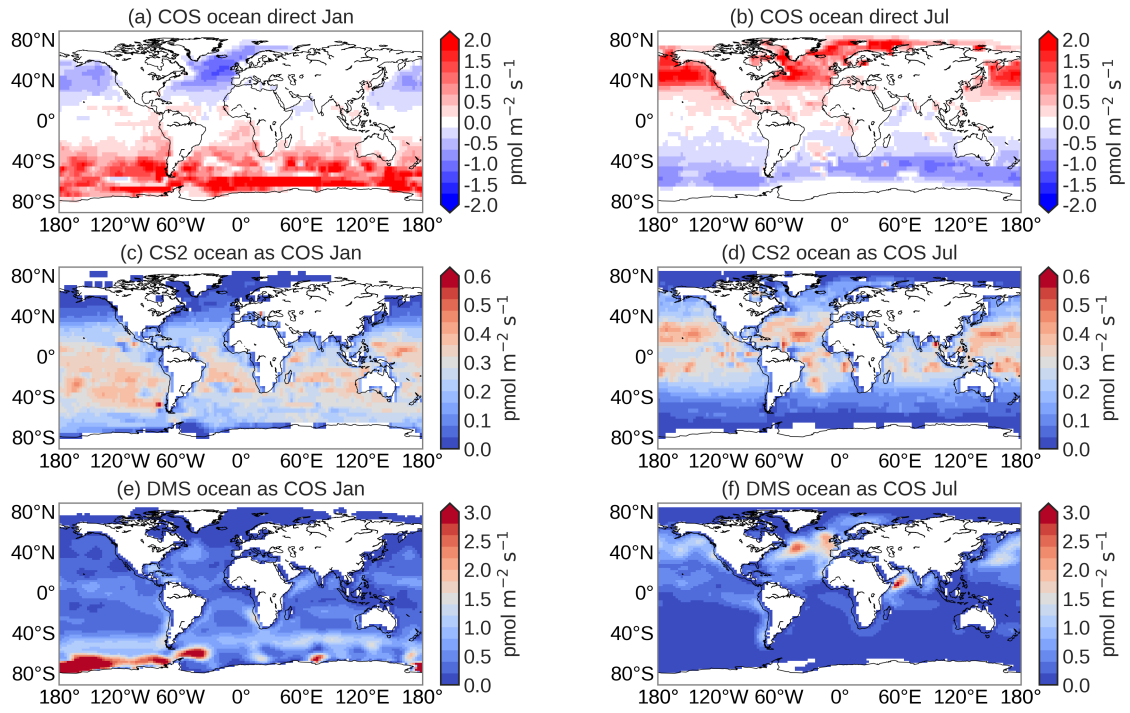


Figure S4. Spatial distribution of the climatological prior ocean fluxes of COS (upper), CS₂ (middle) and DMS (lower) for January (left) and July (right) based on Suntharalingam et al. (2008). Colorbars are extended with an arrow indicating that values may exceed the maximum or minimum value. Fluxes of CS₂ and DMS are converted to COS using yields of 83% and 0.7%, respectively.

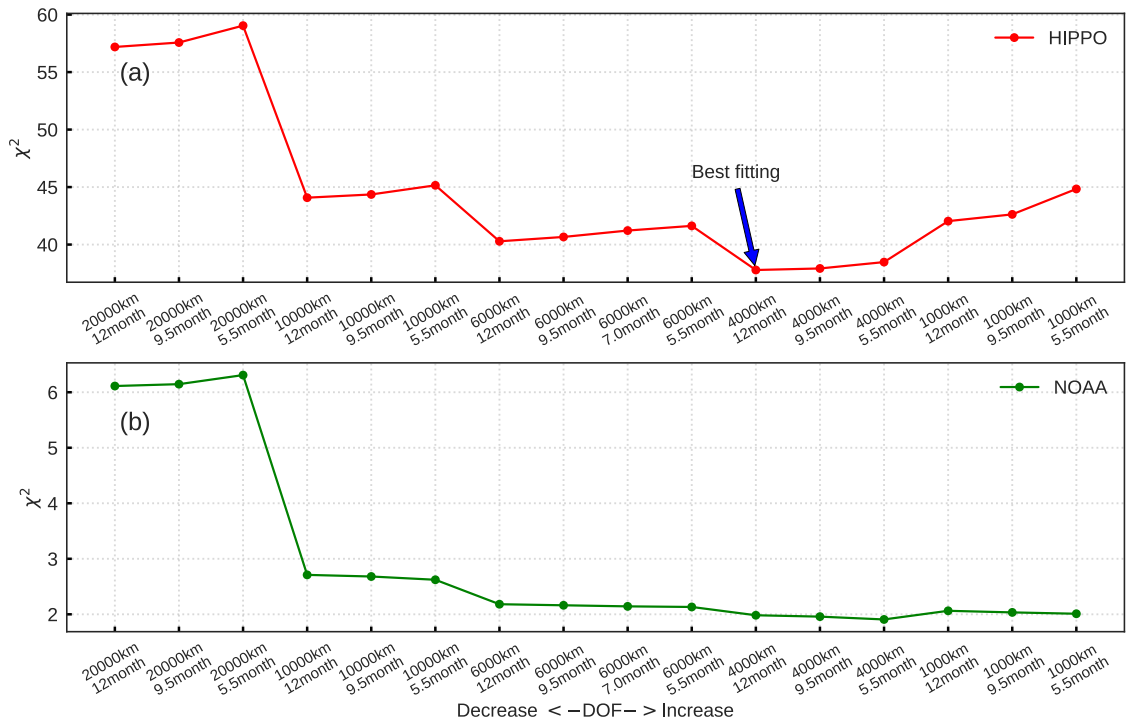


Figure S5. Spatial and temporal correlation settings in inversion S_u and the resulting χ^2 metric. (a) shows the comparison to HIPPO and (b) shows the χ^2 with respect to NOAA surface observations. The inversions were run for the years 2008–2011 and HIPPO campaigns 1-3 were sampled. The first and last 6 months of inversions were removed to calculate χ^2 . The arrow indicates the correlation setting selected for all further experiments.

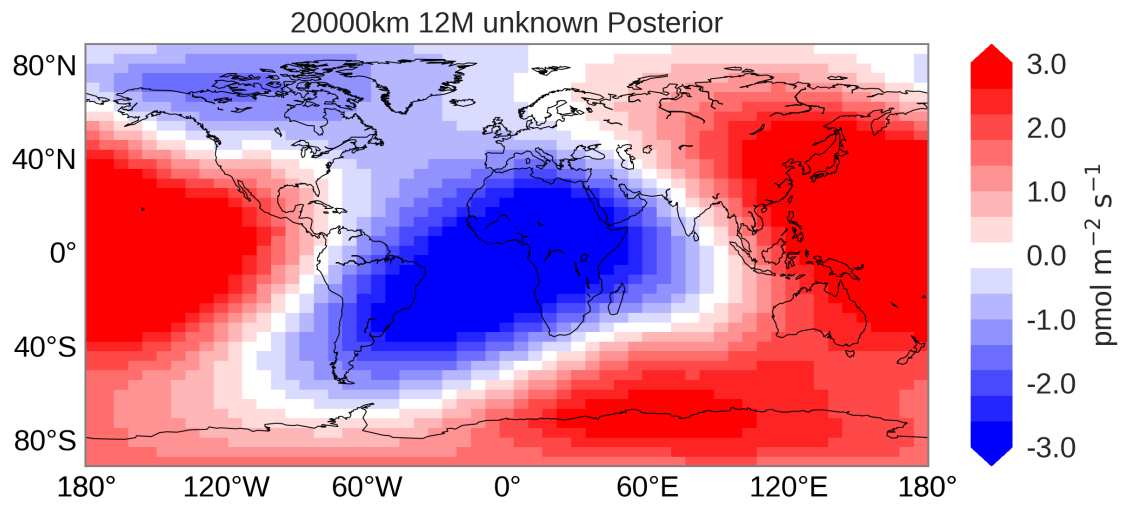


Figure S6. Optimized emission field of inversion S_u for a spatial correlation length of 20000 km and a temporal correlation length of 12 months. This field differs markedly from the optimized patterns presented in the main text.

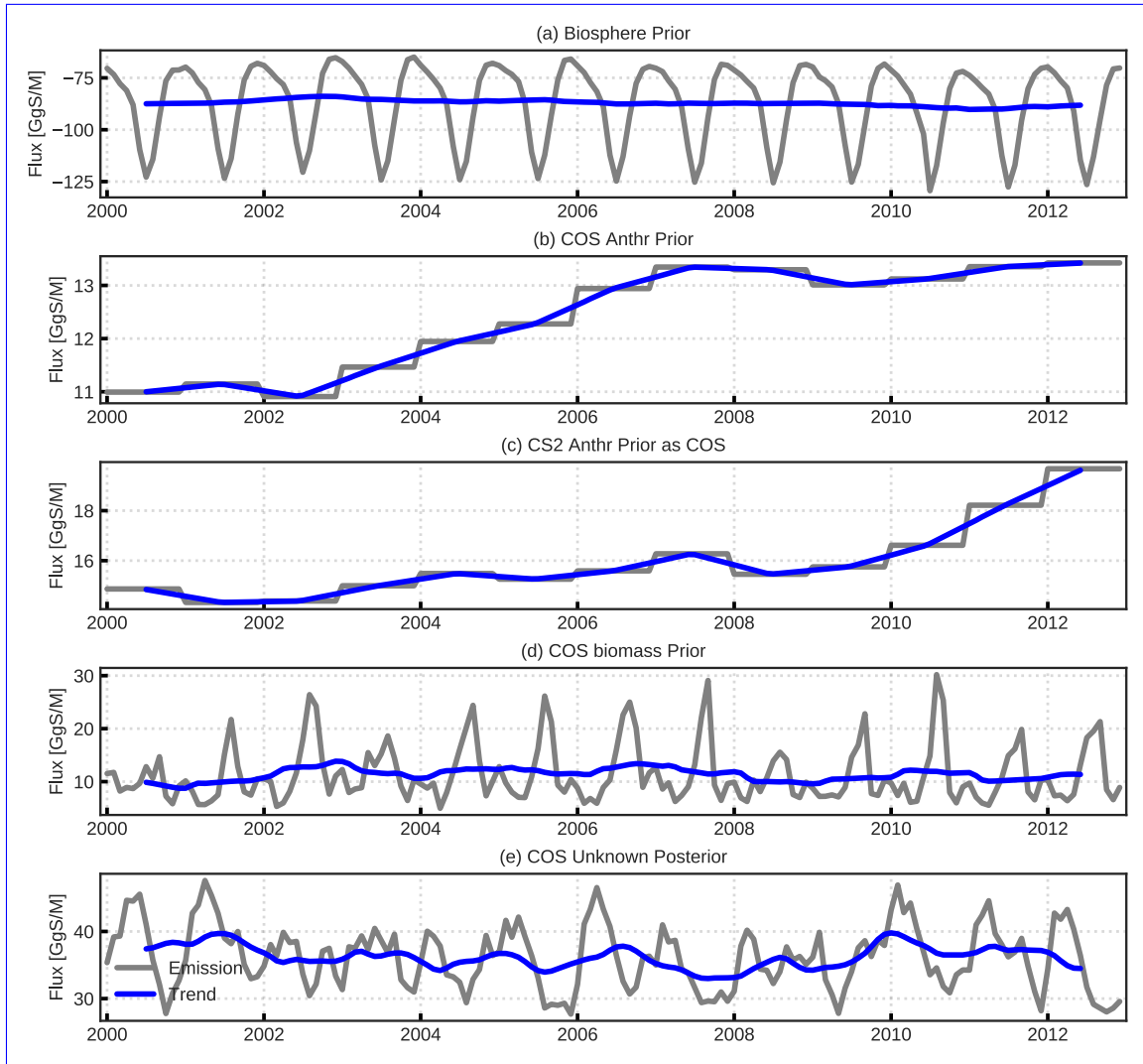


Figure S7. Monthly exchange fluxes related to inversion S_u : (a) prior biosphere (b) anthropogenic COS emissions (c) COS from anthropogenic CS_2 (d) biomass burning (e) optimized "unknown" emissions. Monthly global fluxes are shown in gray, and the trend is shown in blue.

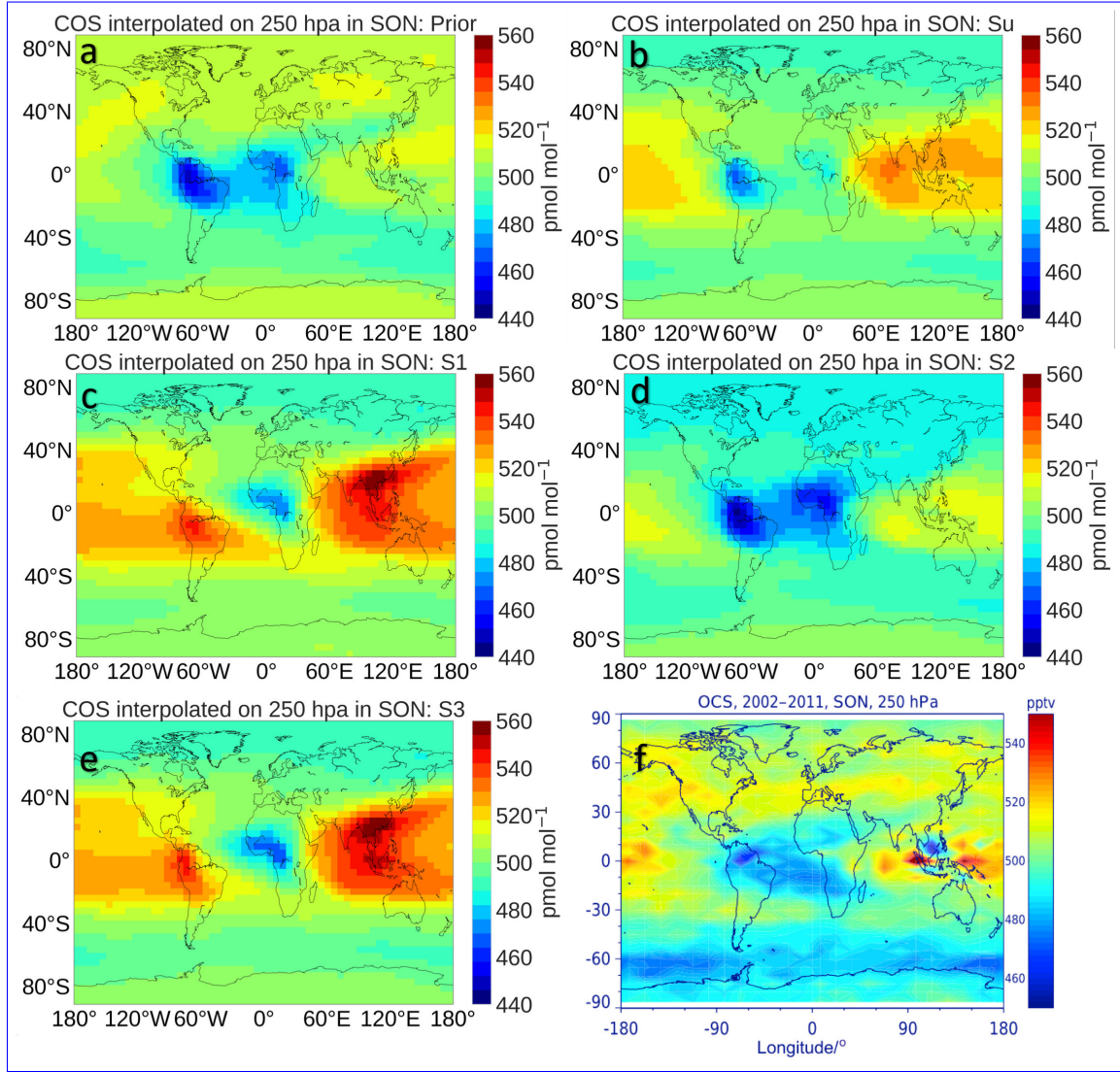


Figure S8. COS mole fraction comparison between MIPAS and TM5 on 250 hPa in September to November. (a–e) are TM5 prior, and inversions S_u , S1, S2, S3, respectively. (f) is captured from figure 11 in (Glatthor et al., 2017). TM5 results represent a 2008–2010 average, and MIPAS is averaged over 2002–2011. Because TM5 is systematically lower than MIPAS, 25 pmol mol⁻¹ is added to the TM5 results for a better visual comparison.

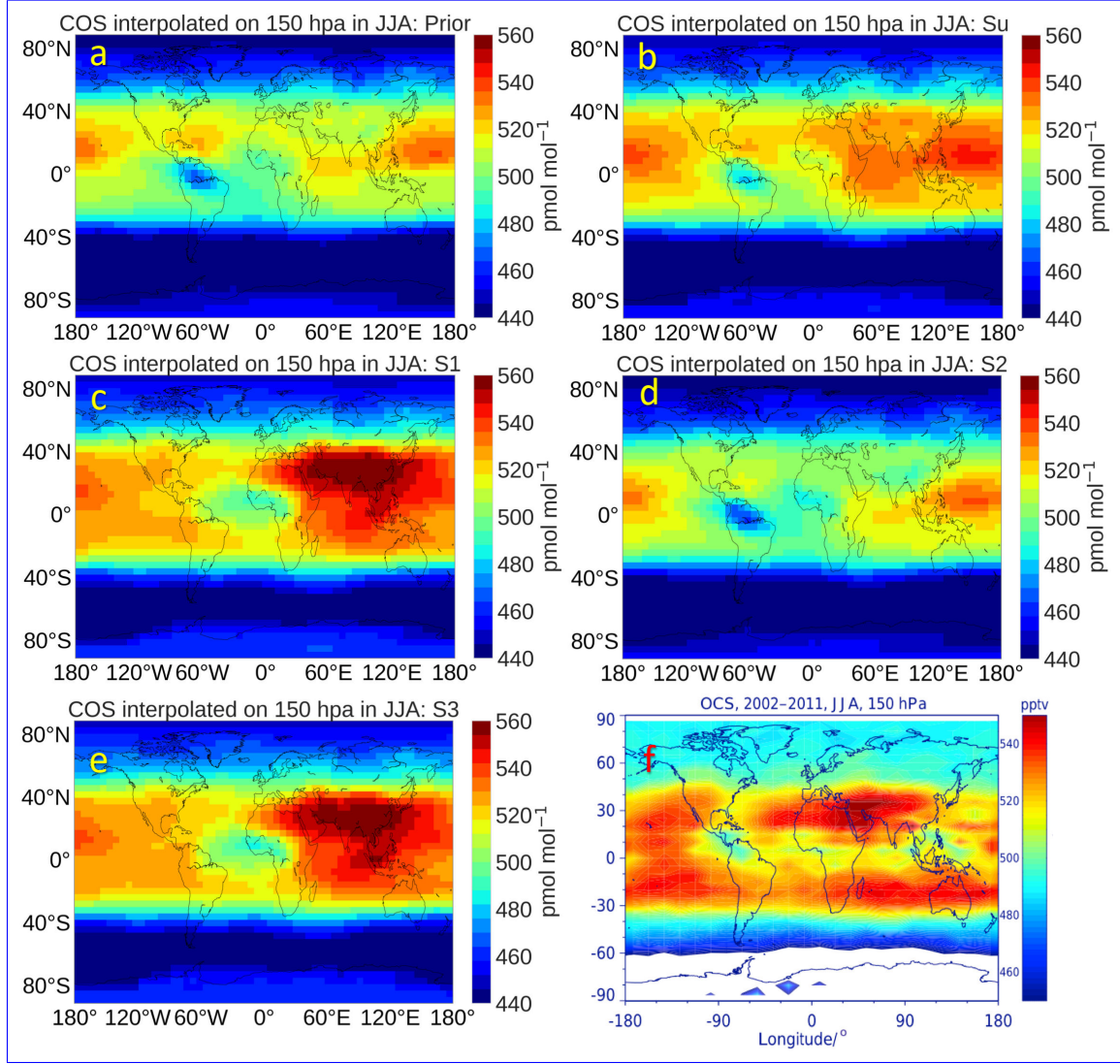


Figure S9. COS mole fraction comparison between MIPAS and TM5 on 150 hPa in June to August. (a–e) are TM5 prior, and inversions S_u , S_1 , S_2 , S_3 , respectively. (f) is captured from figure 12 in (Glatthor et al., 2017). TM5 results represent a 2008–2010 average, and MIPAS is averaged over 2002–2011. Because TM5 is systematically lower than MIPAS, 25 pmol mol^{-1} is added to the TM5 results for a better visual comparison.

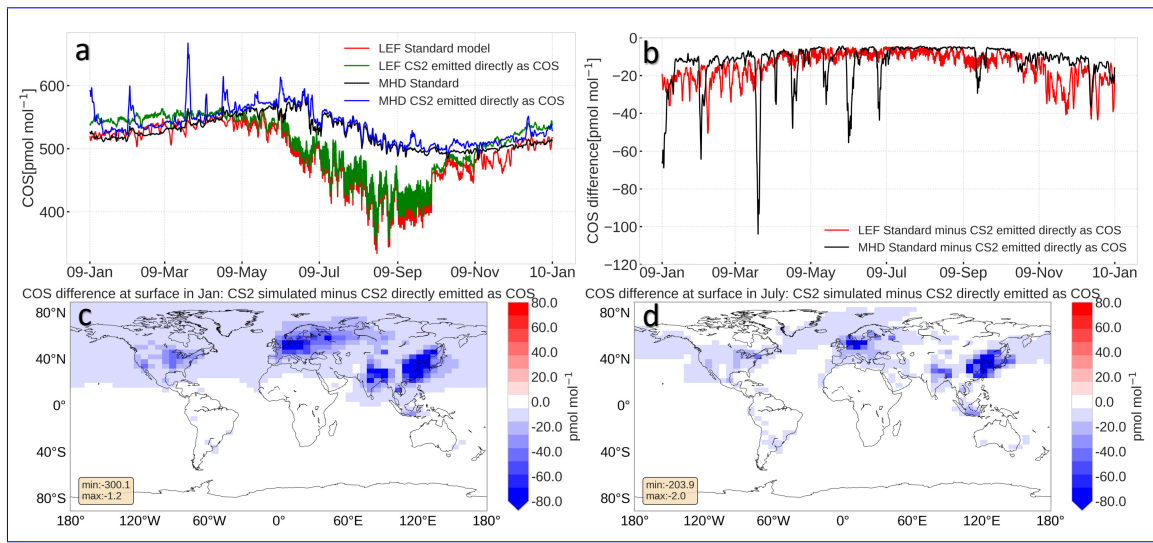


Figure S10. Sensitivity test in which CS₂ is emitted as COS compared to the standard model: (a) COS mole fractions sampled at two stations (LEF and MHD) (b) COS difference sampled at two stations (LEF and MHD) (c) COS spatial difference at the surface in January (d) COS spatial difference at the surface in July. Note that the results are from forward model simulations with prior fluxes in 2008–2010. For clarity, in panels a and b only results in 2009 are shown.

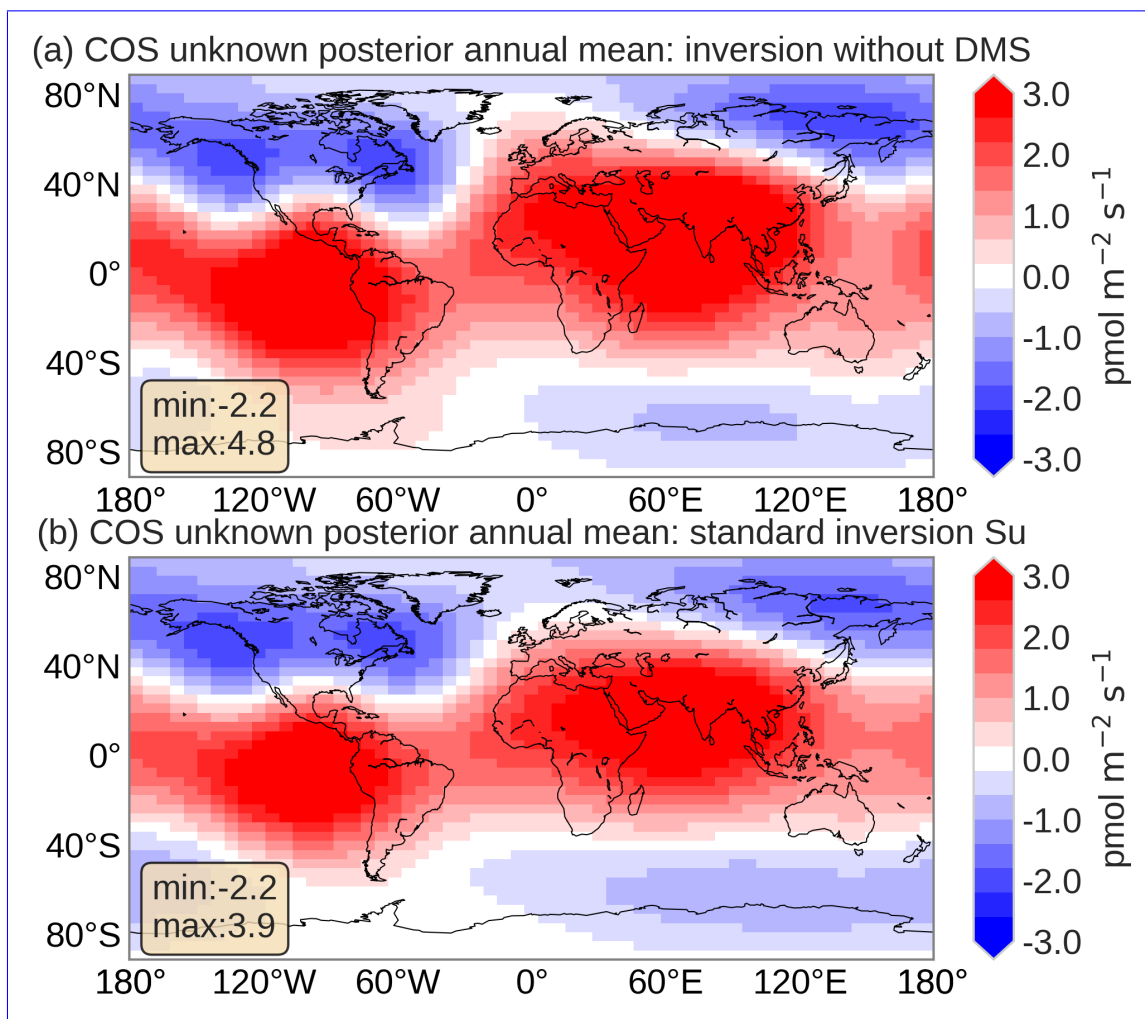


Figure S11. Posterior flux of the NO-DMS inversion and the standard inversion S_u : (a) posterior COS flux with DMS emissions added to the "unknown" flux (b) posterior COS flux from inversion S_u . The inversions have been performed for the year 2008-2010.

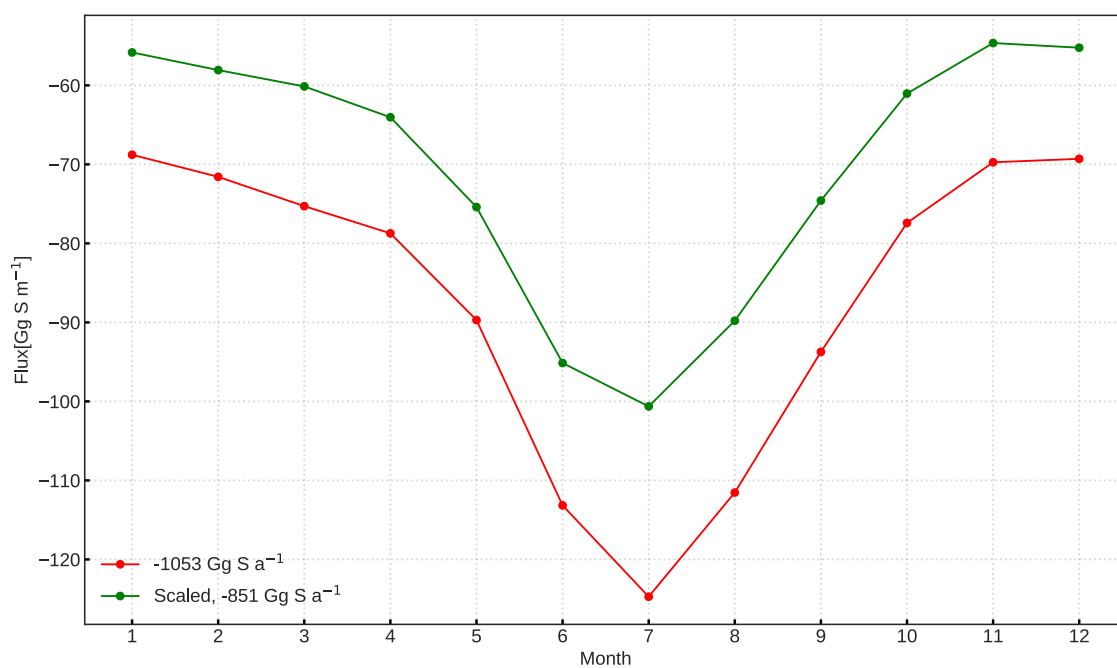


Figure S12. Monthly biosphere flux corrected by scaling with monthly-mean COS mole fractions (green) and the original biosphere flux calculated with a fixed mole fraction of $500 \text{ pmol mol}^{-1}$ used in the paper (red). The yearly-averaged corrected flux amounts to -851 Gg S a^{-1} compared to $-1053 \text{ Gg S a}^{-1}$ for the original flux.

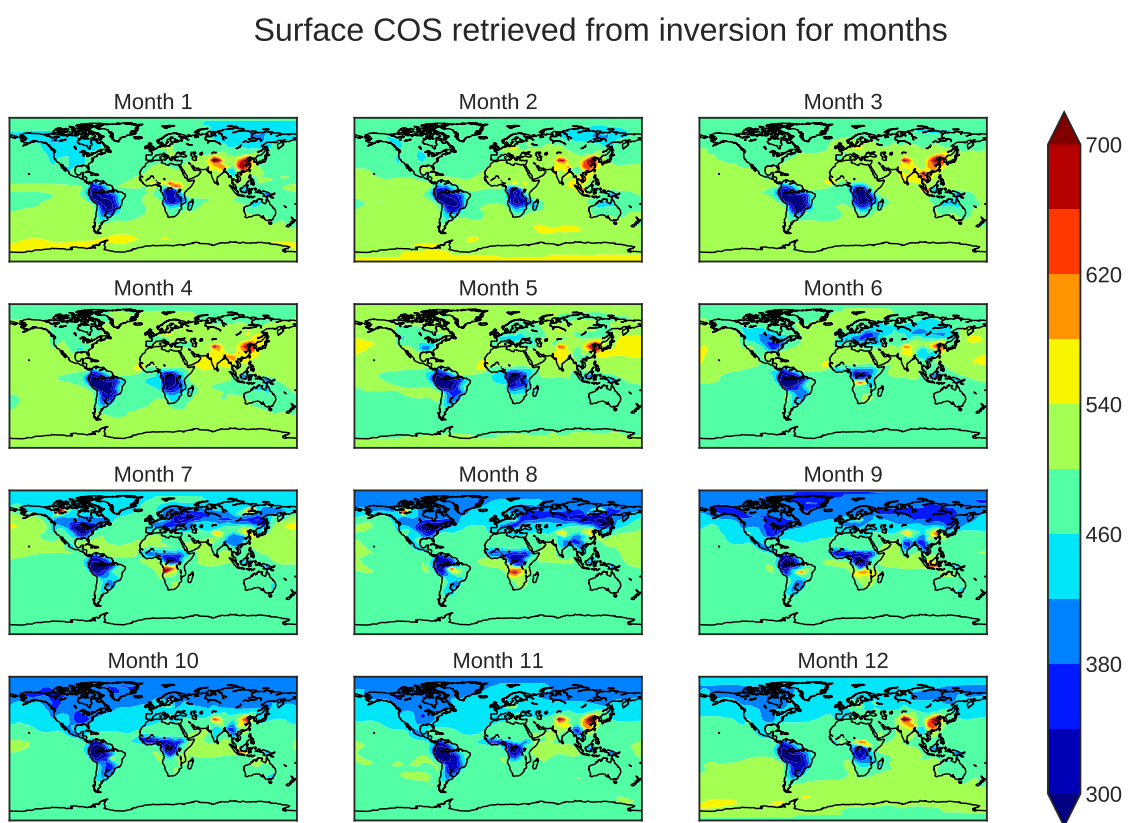


Figure S13. Monthly-mean surface COS mole fractions in pmol mol^{-1} from inversion S_u in the year 2018.

Scaling factor of biosp flux for months

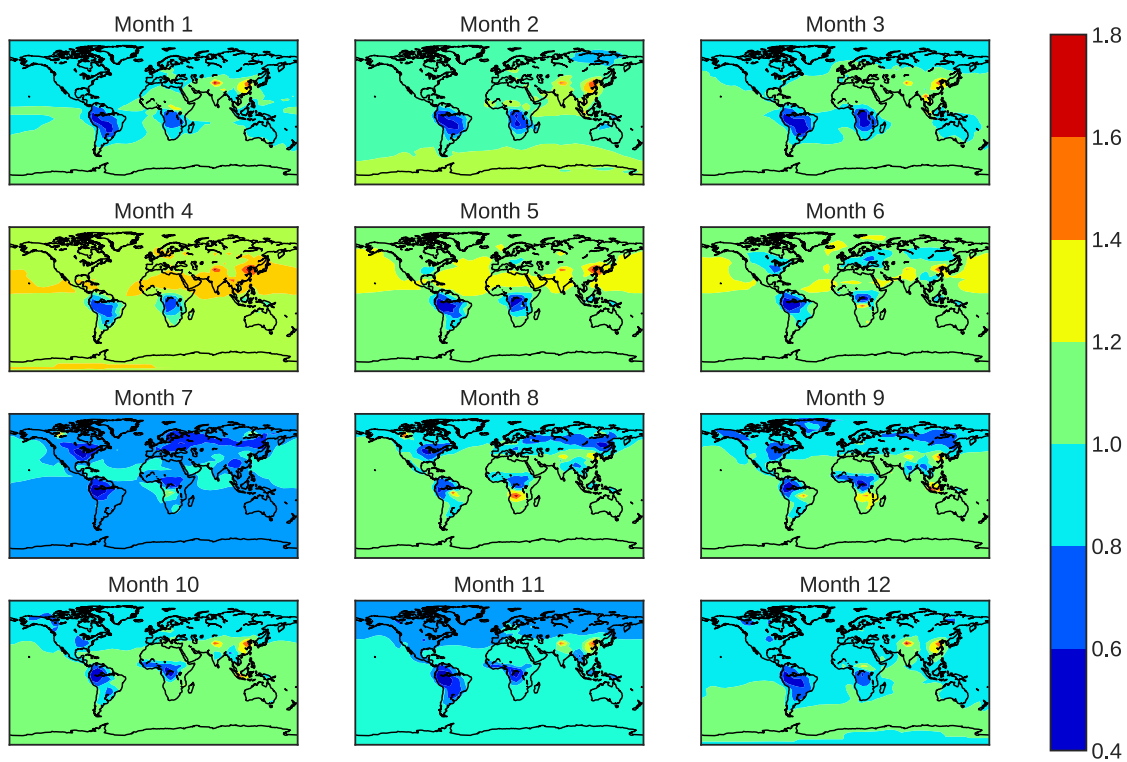


Figure S14. Monthly scaling factors for correcting the original SiB4 prior biosphere flux in the year 2018. The unit-less scaling is calculated as the monthly-mean surface COS mole fraction from Figure S13 divided by $500 \text{ pmol mol}^{-1}$.

Table S1. Prior and posterior fluxes, calculated errors, and error reduction of the different inversions for the 2008–2011 period, aggregated to global totals. Note that we removed the first and last 6 months from the inversions as spin-up and spin-down periods. CS₂ emissions are converted to COS indirect emissions using a conversion factor of 0.83. The correlation lengths for inversions S1–S3 are 4000 km and 12 months.

	Flux category	Grid-scale error	Prior (Gg S a ⁻¹)	Posterior (Gg S a ⁻¹)	Error Reduction
S _u	COS unknown ^a	100 %	432±20.2	436.7±3.8	81.4 %
	COS unknown ^b	100 %	432±25.5	436.7±3.5	86.3 %
	COS unknown ^c	100 %	432±99.0	443.1±7.3	92.6 %
	COS unknown ^d	100 %	432±284.6	423.2±5.2	98.2 %
S1	COS anthropogenic	10 %	157.3±6.9	154.7±6.7	2.4 %
	COS ocean	50 %	40.6±15.4	-15.7±11.8	23.4 %
	COS biosphere	50 %	-1053.1±165.3	-564.0±20.3	87.8 %
	COS biomass	10 %	122.5±4.6	118.7±4.6	0.2 %
	CS ₂ anthropogenic	10 %	190.7±8.1	189.7±8.0	2.1 %
	CS ₂ ocean	50 %	83.0±10.6	98.7±10.5	0.6 %
S2	COS ocean	50 %	40.6±15.4	24.9±6.3	59.0 %
	CS ₂ ocean	50 %	83.0±10.6	535.8±6.5	38.3 %
S3	COS biosphere	50 %	-1053.1±165.3	-613.2±7.1	95.7 %

^a Spatial and temporal correlation length: 1000 km and 5.5 months.

^b Spatial and temporal correlation length: 1000 km and 12 months.

^c Spatial and temporal correlation length: 4000 km and 12 months.

^d Spatial and temporal correlation length: 20000 km and 12 months.

Prior and posterior errors of exchange fluxes are aggregated to global totals and averaged in time by the TM5-4DVAR post-preprocessing (VPP) program, and the statistics are shown in Table S1. The method of error estimation was introduced by Meirink et al. (2008a, b). Because the inverse model system applied in this study is linear, the posterior errors can be estimated, as discussed in Meirink et al. (2008b). Error reduction (ER) is defined as:

$$ER = 1 - \frac{\sigma_{\text{post}}}{\sigma_{\text{prior}}}, \quad (\text{S1})$$

where σ_{post} denotes the posterior error, and σ_{prior} is the prior error. For inversion S_u, only the unknown source term is optimized with a prescribed grid-scale error of 100 %. With increasing spatial and temporal error correlations between the grid-scale prior fluxes, the global prior error increases. As a result, error reductions increase from 81 % (spatial correlation length 1000 km, temporal correlation lengths 5.5 month) to 98 % (20000 km and 12 months), because posterior errors on

10 the global budget closure are all in the range 3–8 %. The error reductions depend strongly on the magnitude of the flux and the error assigned to them. For instance, error reductions for the biosphere fluxes in inversions S1 and S3 are as high as 90 %, which is related to their large magnitude and the large prior error of 50 % assigned to the fluxes. Flux corrections and error reductions of the other S1 fluxes (anthropogenic, biomass burning) are small, indicating that insufficient observations are available to optimize these terms. In inversion S2, in which only oceanic sources were optimized, the error reductions are
15 smaller than for inversions S1 or S3, mostly because ocean fluxes are smaller in magnitude, and give less degrees of freedom to improve the match with observations, especially over land. Note further that some flux terms are optimized way outside the prior error range that was assigned to them. Generally, this indicates that structural model errors are present. In the case of COS, these structural errors are likely related to missing sources and sinks in the COS budget.

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