

Interactive comment on “What are the greenhouse gas observing system requirements for reducing fundamental biogeochemical process uncertainty? Amazon wetland CH₄ emissions as a case study” by A. Anthony Bloom et al.

A. Anthony Bloom et al.

abloom@jpl.nasa.gov

Received and published: 28 October 2016

We thank the reviewers for their constructive feedback and suggested corrections. Below we have addressed each individual comment from reviewers 1 and 2 (reviewer comments are shown in italics; our responses to the reviewer comments are shown in bold). In light of the reviewer comments, our revised manuscript now includes (a) an analysis on the role of CH₄ retrieval systematic biases, and (b) a more robust quantification of the CH₄ flux requirements. We believe that the following revisions have substantially improved the overall quality

C1

of our manuscript.

Anonymous Referee 1

The paper by Bloom et al. investigates the required performance parameters of satellite missions aimed at gaining quantitative insight into the biogeochemical processes driving methane wetland emission in the Amazon region. To this end, the authors first examine the variability (in space, time, magnitude) of the carbon cycle and hydrological processes that control CH₄ emissions. Then, they use observing system experiments to derive mission requirements (spatial and temporal resolution; precision) that would allow for disentangling the processes under natural variability. The study covers satellite concepts in low-earth-orbit (LEO) as well as in geostationary orbit (GEO).

The applied methodology is most interesting since it outlines an approach how to quantitatively derive mission requirements based on the actual variability of the targeted process parameters. I would tend to criticize the study as being too simplistic in one or the other way outlined below. But certainly, the paper is well written, methods are robust and rigorous, and thus, it is suitable for publication in ACP after considering my questions/comments below.

Questions/comments:

(1.1) (1) A shortcoming of the study is the assumption of purely random error sources implying that measurement uncertainty improves with the square root of the number of binned soundings. This assumption results in maps such as Figure 4 where the measurement precision of GEO soundings binned on 300×300 km² is in the range of 0.1 ppb (given 1800 ppb background) which is a clearly unrealistic assumption for the overall measurement error. Experience with the current generation of passive greenhouse gas sounders such as GOSAT and OCO-2 tells that, at aggregated scales, random errors are dwarfed by systematic errors which typically exceed 0.1 ppb by far. Systematic errors are hard to address and, indeed, the manuscript concedes the neglect of systematic errors but a major caveat should be issued when discussing the achievable

C2

flux precisions.

We have now amended our analysis with an explicit simulation of CH₄ residual bias errors. We describe the incorporation of a residual CH₄ bias structure at the end of section 2.3 of the revised manuscript. We now show that the relative advantage of a GEO mission - in comparison to a LEO mission - decreases with increasing CH₄ bias (Figure 7).

(1.2)(2) The manuscript restricts the advantage of a GEO sounder to massively enhanced data density. Wouldn't it make sense to actually exploit the quasi-contiguous temporal sampling of a GEO sounder? A GEO sounder would allow for resolving variability due to source and transport patterns on the time scale of hours. Running an inverse model with monthly flux resolution (and probably imposed sub-monthly variability) might simply discard some of the available process information.

We agree that additional constraints may be achievable under certain process scenarios; for example, emissions from spatially concentrated wetland CH₄ sources (e.g. across the main stem of the Amazon river) could potentially be constrained based on higher resolution CH₄ concentration gradients, and fluxes can be estimated using alternative approaches. Conversely, monthly CH₄ inversions are more suitable for spatially and temporally diffuse CH₄ emission process scenarios. We now discuss the additional potential advantages of GEO OS in the revised manuscript.

Technical comments:

(1.3) P5,L16: Focusing the study on March reduces data amount and related logistics but it neglects seasonal variability. Is there any indication that March is a benign or malign case? For example: is the CH₄ flux precision requirement of 3 mg CH₄/m²/day valid for all seasons?

We have now included a more robust quantification of the CH₄ precision require-

C3

ment: for a given resolution requirement, we derive a year-round precision requirement (now 10 mg m⁻² day⁻¹) as the CH₄ precision needed to statistically distinguish between wetland CH₄ process hypotheses with a 95% confidence.

In the reviewer's words, March 2007 is a "malign case": we now state that "the atmospheric CH₄ OS requirement as the ability to meet the CH₄ flux resolution and precision requirements during the cloudiest time of year". We also clarify that March 2007 is the cloudiest month in the Jan - Apr 2007 season (84% cloud cover) and it is considerably higher than the subsequent dry season (46% - 56% cloud cover).

(1.4) P5,L16: MODIS cannot provide information on diurnal variability in cloud cover. Would you expect a significant effect e.g. for choosing an optimal LEO overpass or for optimizing GEO revisits?

We agree with the reviewer that diurnal variability may amount to a key component of assessing and optimizing GEO and LEO missions. Based on ERA-interim cloud-cover re-analyses, we show that the annual mean diurnal coefficient of variation of cloud-free Amazon basin spans 7% - 80% (median = 29%). Given the non-linear relationship between data yield and 1km x 1km cloud-free domain shown in Figure B1, we highlight that choice of diurnal variability could have a substantial influence on LEO and GEO data yield. We now make these points in the discussion section of our revised manuscript.

(1.5) P7,L8: Looking at the correlation matrix (Figure A1), there is substantial correlation among (C1, C2, C4) and (H1,H2) on spatial scales down to 100 km which means that they would be hard to distinguish by an observing system. So, actually, the requirement $L \leq 300$ km only allows for discriminating carbon and hydrological controls but not for discriminating the type of carbon (except for C3 vs (C1,C2,C4)) or the type hydrological process (except for H3 vs (H1,H2)). Is that correct? Probably, this should be discussed in more detail.

C4

We have now removed this figure from the revised manuscript, since our precision derivation approach implicitly accounts for both spatial and temporal correlations (see response to 1.3).

(1.6) P9,L1: It would be appropriate to cite an original TROPOMI paper at least once (instead of Wecht et al., 2014, repeatedly): P. Veefkind, I. Aben, K. McMullan, H. Forster, J. de Vries, G. Otter, J. Claas, H.J. Eskes, J.F. de Haan, Q. Kleipool, M. van Weele, O. Hasekamp, R. Hoogeveen, J. Landgraf, R. Snel, P. Tol, P. Ingmann, R. Voors, B. Kruizinga, R. Vink, H. Visser, P.F. Levelt, TROPOMI on the ESA Sentinel-5 Precursor: A GMES mission for global observations of the atmospheric composition for climate, air quality and ozone layer applications, Remote Sensing of Environment, Volume 120, 15 May 2012, Pages 70-83, ISSN 0034-4257, <http://dx.doi.org/10.1016/j.rse.2011.09.027>.

We now cite the Veefkind et al., (2012) paper as a reference for the TROPOMI mission.

(1.7) P10,L9: "March and September 2007". The rest of the paper is restricted to March. So, I guess, September needs to be removed.

We have now removed "September".

(1.8) Equation (2): The multiplication of the vectors N and O is not a scalar product but an element-wise multiplication, right? Probably, this needs to be stated somewhere.

We now use an appropriate symbol and explicitly state this in the text.

(1.9) P11,L7: Is the unperturbed CH₄ flux assumed constant (12 mg/m²/day) throughout the domain?

In response to the second reviewer's comments (see responses to 2.2 and 2.9) we now report flux uncertainties in mg m⁻² day⁻¹, and we have revised equation 5 accordingly. Since the explicit definition of f{L,0} is now redundant, it has been removed from the revised manuscript.

C5

(1.10) P11,L17: Figure A2 -> Figure C1

We now correctly reference this figure.

(1.11) P11,L11: A further advantage of GEO is several revisits per day.

We now clearly state this in the revised manuscript

(1.12) Appendices: It would be useful to have a meaningful title for the appendices (instead of only Appenix A, B, C).

We have now added descriptive titles to Appendices A-D.

(1.13) Equation C1: What is the inverse of a vector, f'1?

We have now added a sentence to better clarify that f' is an N x N array, comprised of N flux vectors.

(1.14) P16,L22: Figure A1 -> Figure A2.

Figure reference now corrected

Anonymous Referee 2

This study presents an OSSE for different hypothetical LEO and GEO satellite instruments. The focus is on the requirements on these observing systems for obtaining process-relevant information on wetland emissions in the Amazon region. As explained below some assumptions are made, which are not well justified but have a potentially large influence on the conclusions. These will have to be dealt with in a satisfactory manner to make this paper suitable for publication in ACP.

GENERAL COMMENTS

(2.1) Autocorrelation scales have been derived for several parameters to motivate the choice of spatial scale that the measurements should be able to resolve in order for the OS to help us gain process understanding. It is presented as a novel approach that could be applied to other related problems. Although I appreciate the attempt to de-

C6

rive such scales (which indeed addresses an important question), I do not agree that the presented method solves this problem. The reason is that the results presented in figure 3 depend on the scale of the data sets that are used. What is shown is the autocorrelation of parameters that are averaged on a scale of 0.5x0.5 degree. If the resolution of the datasets were much higher, then other more local processes would contribute to variability shortening the overall auto-correlation scale. Indeed it is questionable whether the methane emission from a local pond really correlates with one that is 100 km away. What is the motivation to use datasets at 0.5x0.5 degree? If the processes themselves motivate this choice then this should be explained. In absence of such a motivation it is a probably more a practical choice. I have no problem with this choice as long as its limitation is made clear, and that it requires reconsideration for any other application.

We agree with the reviewer that our assessment of carbon and hydrological process variable correlation scales requires reconsideration for any subsequent application. We now clarify that the auto-correlation scales are specific to Amazon river basin; we also highlight the limitation of our auto-correlation approach, and we clarify that finer-scale analyses may require higher resolution datasets to quantify GHG measurement requirements.

We also agree with the reviewer that finer-scale variability from higher-resolution datasets could potentially contribute to alternative assessments of auto-correlation scales. However, in our derivation of Moran's I at each L, we aggregate our data at an L x L resolution (see Appendix A), and therefore fine-scale variability is averaged out (regardless of the native resolution of the dataset).

(2.2) If it is considered important that the inversion resolves the autocorrelation scale then it is not sufficient to evaluate the posterior uncertainty at that scale. This is because the off-diagonals of the posterior covariance matrix might indicate that neighboring fluxes are not independently determined. In this study, however, the performance criterion only considers values on the diagonal. In addition, the choice of 25% confuses

C7

monthly and annual fluxes. The requirement is on monthly fluxes, but it is derived from an estimate of Melack et al on the annual time scale.

We agree with the reviewer that using a “%” precision is misleading. We now present flux precision in CH₄ flux units (mg m⁻² day⁻¹) throughout the manuscript; the units are now consistent with our revised precision requirement (10 mg m⁻² day⁻¹; see response to reviewer comment 1.3).

We agree with the reviewer that “off-diagonal” error correlations in retrieved fluxes would likely indicate that neighbouring fluxes are not independently determined. However, as long as all diagonal terms meet the precision requirement (10 mg m⁻² day⁻¹), the OS can resolve underlying spatial flux patterns at the required precision (regardless of posterior error covariance).

(2.3) It is unclear why a special effort is made to derive requirements on horizontal resolution looking at the drivers of processes, whereas this is not done for the requirements on flux precision and temporal resolution. Since the inversion solves for net fluxes it remains unclear anyway if these requirements really allow us to constrain specific processes. Wouldn't it have been more logical to vary process model parameters to determine what is needed to resolve them? You might wonder whether it is even realistic to constrain processes only by measuring XCH₄ using a single instrument. Atmospheric measurements are useful for constraining regional emission budgets, which - in combination with other information - can be used to derive improved process understanding. The OSSE approach that is taken disqualifies instruments that provide useful constraints on larger scales as part of a multi-component global monitoring system.

We now include a quantification of the CH₄ flux precision requirements for distinguishing between both spatial and temporal CH₄ emission hypotheses (see response to reviewer comment 1.3). We have also now included a lagged Pearson's correlation analysis to determine the temporal process control correlation lengths.

C8

We agree with the reviewer that varying process parameters in a model is potentially a useful approach for quantifying the OS needed to improve process understanding. However, due to the scarcity of top-down constraints and in-situ measurements in tropical wetland environments, little is known about whether current models are able to capture the first-order spatial and temporal variability of wetlands. We discuss and state that model approaches can be used - albeit with due caution - to define CH₄ OS measurement requirements in of the revised manuscript.

Finally, we now highlight the need to investigate the added advantages of a multi-component global monitoring system in the revised manuscript.

(2.4) This OSSE is extremely (and unrealistically I would say) optimistic about the uncertainty reduction that can be achieved by averaging large numbers of data. It is mentioned that the 'cumulative' uncertainty of GEO OS may be as low as 0.02 ppb. It is probably a main reason why the GEO measurement concept performs so well in this study. In reality, however, systematic uncertainties will kick in at much reduced precisions preventing any further improvements upon averaging. Some attempt should be made to assess the sensitivity of the conclusion that improved process-understanding calls for the GEO approach, to the presence of systematic errors in the data.

We agree with the reviewer that systematic biases are a limiting factor in the potential performance of a GEO approach. We have now included a residual CH₄ bias analysis to address this comment (see response to comment 1.1).

(2.5) Further effort is needed to quantify the impact of errors due to the simplified treatment of atmospheric transport. In general, surface fluxes are proportional to spatio-temporal concentration gradients in the atmosphere. Looking at figure C1 it becomes clear that the east-west gradient in WRF is substantially stronger than in LPDM. It has probably to do with the north- and southward transport along the Andes in WRF, which is missing in LPDM. The impact of this should be quantified.

C9

We agree with the reviewer that the LPDM approach underestimates the east-west gradient (see response to 2.18), and we now highlight that the LPDM provides a conservative estimate on the observable CH₄ gradients across the region. To quantify the potential bias stemming from underestimated CH₄ gradient across the Amazon domain, we conduct a sensitivity test on the GEO and LEO median flux precision retrievals, where the LPDM-derived transport operator is multiplied by 1.5. We find that this leads to an inversely proportional (33%) reduction in the GEO and LEO flux precision (the sensitivity test results are reported in the revised manuscript).

(2.6) It should be made clearer why the analysis is limited to the month of March. Many things are different in other months (atmospheric dynamics, cloud cover, CH₄ fluxes, etc.). March doesn't sound like a particularly good choice as average, or representative month.

We now define our OS requirements as the ability to resolve monthly CH₄ fluxes at the required resolution and precision during the cloudiest part of the 2007 wet season (see response to comment 1.3). We also highlight that March is the cloudiest month in the 2007 wet season. Finally, we highlight the need to investigate the role seasonal transport variability (amongst other factors) on GEO and LEO CH₄ flux retrievals.

SPECIFIC COMMENTS

(2.7) Page 7, line 14: "Throughout ... CH₄ emissions" I don't see why the fact that 25% is in between the dynamic ranges of monthly GPP and inundation variability would make it suitable for separating their influences. Apart from this, what justifies the assumed linearity between these drivers and methane emissions?

We have now addressed this concern with a more robust derivation of CH₄ flux requirements (see response to comment 1.3).

C10

(2.8) Page 9, line 11: 'i.e. all accepted ... 100% cloud-free' According to Appendix B, MODIS data that is probably cloud-free are considered as fully cloud-free. These two statements do not fit together.

We have grouped “probably cloud free” and “cloud free” flags together, and “probably cloudy” and “cloudy” flags together. We have clarified this in the revised manuscript, and we have added a sentence in the appendix to clarify our assumption.

(2.9) Page 11, equation 3: Why is $c\{L,0\}$ calculated? In the end all that matters is the spread in 'c' due to the random perturbation and how it maps on 'f' using 'A'. The uncertainty in 'f' does not depend on the mean of 'c'.

We agree with the reviewer's statement, since our derivation of f (equation 5) is independent of $c\{L,0\}$. For the sake of simplicity, we now set all $c\{L,0\}$ values to zero.

(2.10) Page 13, line 21: 'If Amazon CH_4 fluxes likely be lower' This depends on the distribution of cloud cover. The wettest regions will likely be measured the least frequent. This calls for further motivation of why uniform emissions have been assumed.

In the revised manuscript, we now clearly define our OS requirement as the ability to statistically distinguish between biogeochemical process hypotheses based on cloud cover statistics during the cloudiest time of the 2007 wet season (see response to comment 1.3).

(2.11) Page 15, line 7: Why is the purpose of the parentheses here? Please clarify further at what p-level the autocorrelations are required to be significant, and how this is determined. For example in the following sentence it is not clear what r_i refers to. Please revise the description to explain more clearly what was done.

We have now revised this sentence to better convey our derivation of the Moran's

C11

I p-value.

(2.12) Figure 1: What are the different lines in the inset figure?

The green lines denote the average WETCHIMP model Amazon basin monthly CH_4 emissions. We have revised the figure caption to clarify this.

(2.13) Figure 4: Why do you call this 'cumulative precision'? Isn't it rather the precision of a $300 \times 300 \text{ km}^2$ average?

We now explicitly define CH_4 “cumulative precision” in the revised manuscript. For the sake of clarity, we also define CH_4 “cumulative precision” in the figure caption.

(2.14) Figure 5: Why isn't cloud filtering affecting the number of data, comparing GEO, GEO-Z1, GEO-Z2?

Observations per unit area include all attempted measurements (both cloud and cloud-free measurements). We have revised the figure caption to reflect this.

(2.15) Figure B1: I assume that both panels represent March 2007. If so, then this should be made clear.

Figure caption updated

(2.16) Figure C1: Do these values represent the total column? If so, then mention this.

Figure caption updated

(2.17) Appendix B, line 18: $f(\omega, i)$ is not used in equation 1. Where do the $30 \times 30 \text{ km}^2$ areas come from?

We now correctly use 'phi' (as opposed to 'f') in referencing the fraction of cloud-free observations in equation 1. We have also corrected ' $30 \times 30 \text{ km}^2$ ' to ' $L \times L$ '.

(2.18) Appendix C, line 17: The mean in CH_4 is not the relevant quantity to compare

C12

LPDM and WRF (it is the gradient in the wind direction that matters).

We now also report the LPDM-approach and WRF gradients across the domain in Appendix D (13.14ppb and 17.24ppb respectively); we calculate the gradients as the CH₄ difference between the North-East and South-West sub-regions of Amazon basin domain. We have also updated the LPDM-WRF figure to mark the delineation between the “North-East” and “South-West” regions.

Interactive comment on Atmos. Chem. Phys. Discuss., doi:10.5194/acp-2016-325, 2016.