

Constraints on methane emissions in North America from future geostationary remote sensing measurements

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Abstract. The success of future geostationary (GEO) satellite observation missions depends on our ability to design instruments that address their key scientific objectives. In this study, an Observation System Simulation Experiment (OSSE) is performed to quantify the constraints on methane (CH₄) emissions in North America obtained from Short Wave Infrared (SWIR), Thermal Infrared (TIR),
5 and multi-spectral (SWIR+TIR) measurements in geostationary orbit and from existing SWIR low earth (LEO) measurements. An efficient stochastic algorithm is used to compute the information content of the inverted emissions at high spatial resolution ($0.5^\circ \times 0.7^\circ$) in a variational framework using the GEOS-Chem chemical transport model and its adjoint. Our results show that at sub-weekly time-scales, SWIR measurements in GEO orbit can constrain about twice as many independent flux
10 patterns than in LEO orbit, with a degree of freedom for signal (DOF) for the inversion of 266 and 115, respectively. Comparisons between TIR GEO and SWIR LEO configurations reveal that poor boundary layer sensitivities for the TIR measurements cannot be compensated by the high spatiotemporal sampling of a GEO orbit. The benefit of a multi-spectral instrument compared to current SWIR products in a GEO context is shown for sub-weekly time-scale constraints, with an increase in the
15 DOF of about 50% for a 3-day inversion. Our results further suggest that both the SWIR and multi-spectral measurements on GEO orbits could almost fully resolve CH₄ fluxes at a spatial resolution of at least $100\text{km} \times 100\text{km}$ over source hotspots (emissions $> 4 \times 10^5 \text{ kg day}^{-1}$). The sensitivity of the optimized emission scaling factors to typical errors in boundary and initial conditions can reach 30% and 50% for the SWIR GEO or SWIR LEO configurations, respectively, while it is smaller than
20 5% in the case of a multi-spectral GEO system. Overall, our results demonstrate that multi-spectral measurements from a geostationary satellite platform would address the need for higher spatiotem-

poral constraints on CH₄ emissions while greatly mitigating the impact of inherent uncertainties in source inversion methods on the inferred fluxes.

1 Introduction

25 Methane (CH₄) plays a key role in both atmospheric chemistry composition and climate. With a radiative forcing relative to preindustrial times that is one third that of carbon dioxide, CH₄ is the second most important greenhouse gas (Myhre, 2013). Further, as a precursor to tropospheric ozone, CH₄ also impacts surface-level air quality (Fiore et al., 2002; West et al., 2006; West and Fiore, 2005), crops (e.g., Shindell et al., 2012), and contributes to ozone radiative forcing (e.g., Fiore et al., 30 2008). Considerable uncertainty remains in our understanding of CH₄ sources (e.g., Dlugokencky et al., 2011; Kirschke et al., 2013), which include emissions from coal, wetlands, livestock, landfills, biomass burning, geologic seepage, and leaks from the production and distribution of natural gas.

Although there is a growing interest in using CH₄ emission regulations as an efficient lever to simultaneously address current air quality and global warming challenges (e.g., West et al., 2012), 35 the lack of confidence in the available CH₄ emission estimates remains a problematic limitation to design of efficient environmental policies. Indeed, recent studies showed discrepancies of up to a factor of two between bottom-up inventories and top-down inversions using atmospheric CH₄ concentration observations (Zavala-Araiza et al., 2015; Katzenstein et al., 2003; Kort et al., 2008; Xiao et al., 2008; Karion et al., 2013; Miller et al., 2013; Wecht et al., 2012; Caulton et al., 2014; 40 Turner et al., 2015; Wecht et al., 2014a). Extrapolation of local emission characteristics to larger areas and/or the use of proxy data (e.g., energy consumption, emission ratios applied to co-emitted species) are the main sources of error in bottom-up methods (Zhen et al., 2015). On the other hand, top-down approaches using space-based measurements of CH₄ from Low Earth Orbit (LEO) platforms allow a global spatial coverage within one to six days but at the same local time. However, as 45 CH₄ emissions can exhibit significant diurnal cycles, e.g., over wetland or boreal peatland (Morin et al., 2014; Gazovic et al., 2010), such temporal undersampling may affect our ability to accurately quantify those fluxes. More generally, insufficient observational coverage and the diffusive nature of transport considerably reduce our ability to spatially resolve grid-scale emissions from space.

Geostationary (GEO) remote-sensing measurements would alleviate the above mentioned shortcomings by providing an almost continuous monitoring and complete spatial coverage of CH₄ concentrations within the field of view. Previous studies have already demonstrated the potential of column-integrated trace gas measurements from geostationary satellites to constrain surface fluxes at regional scale, from single mega-city emissions down to power plant sources (Polonsky et al., 2014; Rayner et al., 2014). The GEOstationary Coastal and Air Pollution Events (GEO-CAPE) mission 55 (Fishman et al., 2012) was recommended by the National Research Council's Earth Science Decadal Survey in order to improve our understanding of both coastal ecosystems and air-quality from re-

gional to continental scale. Its aim is to enable multiple daily observations of key atmospheric and oceanic constituents over North and South America from a GEO platform. For air-quality applications, such high-spatial and high-temporal-resolution measurements would enable source estimates of air quality pollutants and climate forcers and development of effective emission-control strategies at an unprecedented level of confidence. In order to provide more flexibility and to minimize the cost and risk of the mission, the concept of a phased implementation that would launch remote-sensing instruments separately on commercial host spacecrafts has been adopted. The first phase will consist of the launching of the Tropospheric Emissions: Monitoring of Pollution (TEMPO) instrument circa 2019 (Chance et al., 2013), which will provide GEO hourly measurements of ozone and precursors as well as aerosols over greater North America (from Mexico City to the Canadian tar sands, and from the Atlantic to the Pacific ocean). For the second phase, which aims at completing GEO-CAPE's mission requirements by including measurements of important drivers of climate and air quality such as CH₄, CO, and ammonia (Zhu et al., 2015), a rigorous instrument design study is critical to achieve the mission's scientific objectives within its budget constraints.

In this study we perform an Observation System Simulation Experiment (OSSE) in order to characterize the constraints on grid-scale CH₄ emissions over North America provided by different potential GEO-CAPE instrument configurations. The simulation consists of a 4D-Var inversion of CH₄ emissions using the GEOS-Chem chemical-transport model (CTM) over a $0.5^\circ \times 0.7^\circ$ horizontal grid resolution covering North America. In practice, quantifying the information content of such a high-dimensional problem requires either Monte-Carlo simulations or, for linear models, a numerical approximation of the inverse Hessian matrix of the 4D-Var cost function (Tarantola, 2005). *Although previous studies have used Monte-Carlo estimates (e.g., Chevallier et al., 2007; Liu et al., 2014; Cressot et al., 2014), their computational cost can be prohibitive, since many perturbed inversions (typically about 50) are needed, each of them usually requiring numerous forward and adjoint model integrations. Alternatively, inverse Hessian approximations based on information from the minimization itself can be employed, but are usually of very low rank (e.g., Meirink et al., 2008; Bousserez et al., 2015). Therefore, most information content analyses in previous trace-gas Bayesian inversion studies have relied on explicit calculations of the inverse Hessian matrix, by either considering a regional domain (e.g., Wecht et al., 2014a) or performing a prior dimension reduction of the control vector (e.g., Wecht et al., 2014b; Turner and Jacob, 2015). However, thus far dimension reduction methods for high-dimensional problems have relied on suboptimal choices for the reduced space, which preclude an accurate and objective quantification of the spatio-temporal constraints on the optimized emissions.*

In this study we use a gradient-based randomization algorithm to approximate the inverse Hessian of the cost function (Bousserez et al., 2015), which allows us to calculate the posterior errors as well as the model resolution matrix (or averaging kernel) of our CH₄ emission inversion at grid-scale resolution. Such information is used to evaluate the impact of different instrumental designs

(spatio-temporal sampling, vertical sensitivity of the measurements) on CH₄ emission constraints.

95 In particular, the potential of CH₄ retrievals from existing Short Wave Infrared (SWIR) and Thermal
 Infrared (TIR) measurements as well as from a hypothetical multi-spectral instrument on geostation-
 ary orbit are examined. Section 2 describes the Observing System Simulation Experiment (OSSE)
 framework considered in this study, which comprises the 4D-Var method, the forward model, as
 well as the observations and prior information used. Section 3 presents the results of our experi-
 100 ments, where the information content of the inversion is analyzed in detail. A conclusion to this
 work is presented in the last section of the paper.

2 Inverse method

2.1 4D-Var system and information content

The variational approach to Bayesian inference is the method of choice for high-dimensional prob-
 105 lems, since the solution can be computed by iteratively minimizing a cost function instead of alge-
 braically solving for the minimum, which becomes computationally intractable for high-dimensional
 systems. Provided the error statistics are all Gaussian, finding the maximum likelihood entails solv-
 ing the following problem:

$$\arg \min_{\mathbf{x}} J(\mathbf{x}) \quad (1)$$

$$110 \quad J(\mathbf{x}) = \frac{1}{2} (\mathbf{H}(\mathbf{x}) - \mathbf{y})^T \mathbf{R}^{-1} (\mathbf{H}(\mathbf{x}) - \mathbf{y}) + \frac{1}{2} (\mathbf{x} - \mathbf{x}_b)^T \mathbf{B}^{-1} (\mathbf{x} - \mathbf{x}_b),$$

where \mathbf{x}_b is the prior vector, defined in the control space E of dimension n , \mathbf{x} belongs to E , \mathbf{y} is
 the observation vector, defined in the observations vector space F of dimension p , $H : E \rightarrow F$ is
 the forward model operator (also called observational operator), which associates to any vector in
 115 E its corresponding observation in F , and \mathbf{R} and \mathbf{B} are the covariance matrices of the observation
 and prior errors with dimension $(p \times p)$ and $(n \times n)$, respectively. The argument of the minimum of
 Eq. (1) is called the analysis and is referred to as \mathbf{x}_a .

When the adjoint of the forward model (\mathbf{H}^T) is available, the minimum of the cost function J can
 be found iteratively using a gradient-based minimization algorithm (Lions, 1971). The gradient of
 120 the cost function with respect to the control vector \mathbf{x} can be written:

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

An important result is that if the forward model is approximately linear the posterior error covari-
 ance matrix \mathbf{P}^a is equal to the inverse of the Hessian of the cost function:

$$125 \quad \mathbf{P}^a = (\nabla^2 J)^{-1}(\mathbf{x}_a) = (\mathbf{B}^{-1} + \mathbf{H}^T \mathbf{R}^{-1} \mathbf{H})^{-1}. \quad (3)$$

This equivalence can be used to compute information content diagnostics prior to performing
 the inversion. In this study, following Bousserez et al. (2015), the diagonal elements of \mathbf{P}^a (error

variances) are computed using a randomization estimate of $\mathbf{H}^T \mathbf{R}^{-1} \mathbf{H}$. Here an ensemble of 500
 130 random gradients of the cost function are used, based on the convergence of the uniform norm ($\|\cdot\|_\infty$)
 of the inverse Hessian approximation. Bousserez et al. (2015) showed that good approximation of
 both the error variances and the error correlations can be obtained using this approach. For the
 present study we further validated our method by comparing direct finite-difference estimates of
 selected diagonal elements of \mathbf{P}^a to their stochastic approximations and found a relative error smaller
 135 than 10 %.

The model resolution matrix (or averaging kernel \mathbf{A}) is defined as the sensitivity of the analysis
 \mathbf{x}_a (optimized CH_4 emissions) to the truth \mathbf{x}_t (true emissions):

$$\mathbf{A} \equiv \frac{\partial \mathbf{x}_a}{\partial \mathbf{x}_t}. \quad (4)$$

140 The model resolution matrix in Eq. (4) can be rewritten in matrix form:

$$\mathbf{A} = \mathbf{I} - \mathbf{P}^a \mathbf{B}^{-1}. \quad (5)$$

Since \mathbf{B} is diagonal in our experiments, Eq. (5) allows us to calculate any element of \mathbf{A} using:

$$\mathbf{A}_{i,j} = \delta_{ij} - \frac{\mathbf{P}_{i,j}^a}{\mathbf{B}_{j,j}}. \quad (6)$$

145 Finally, the degree of freedom for signal (DOF) of the inversion is defined as the trace of \mathbf{A} , that is:
 $\text{DOF} = \sum_i \mathbf{A}_{i,i}$.

2.2 Forward model and prior emissions

The forward model in Eq. (1) includes the GEOS-Chem chemistry-transport model, which relates
 150 the CH_4 emissions to the 3-D concentration field of atmospheric CH_4 , and the satellite observation
 operator that transforms the CH_4 concentration profiles into their corresponding retrieved profile or
 columns. The GEOS-Chem simulation used in our experiment is described in Wecht et al. (2014a);
 Turner et al. (2015). It consists of a nested simulation over North America at $0.5^\circ \times 0.7^\circ$ horizontal
 resolution and 72 vertical levels, driven by offline meteorological data provided by GEOS-5 reanaly-
 155 sis from the NASA Global Modeling and Assimilation Office (GMAO). Boundary conditions for the
 nested domain are used every three hours from a global $4^\circ \times 5^\circ$ GEOS-Chem simulation. In the case
 of profile assimilation (multi-spectral instrument), the application of the measurement averaging
 kernels to the model profiles can be written as follows:

$$\ln \mathbf{z}_{\text{retr}} = \ln \mathbf{z}_a + \mathbf{A}(\ln \mathbf{z}_{\text{mod}} - \ln \mathbf{z}_a), \quad (7)$$

where \mathbf{z}_{retr} is the profile that would be retrieved if the modeled profile concentrations (\mathbf{z}_{mod}) were
 sounded, and \mathbf{z}_a represents the prior profile concentrations. In the case of X_{CH_4} columns assimi-
 lation, we obtain (Parker et al., 2011):

$$\text{X}_{\text{CH}_4} = \frac{\text{X}_{\text{CO}_2}}{\Omega_{\text{CO}_2}} (\Omega_a + \mathbf{a}^T (\boldsymbol{\omega}_{\text{mod}} - \boldsymbol{\omega}_a)), \quad (8)$$

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where ω_{mod} is the modeled vertical profile of methane, ω_a is the a priori profile, Ω_a is the corresponding a priori column concentration of methane, \mathbf{a} is a column averaging kernel vector that describes the sensitivity as a function of altitude, ω is the measured vertical column concentration of CO_2 , and X_{CO_2} is a modeled column mixing ratio of CO_2 . For simplicity, we use a single averaging kernel for each instrument. A larger ensemble of averaging kernels describing a potential range of sensitivities is beyond the scope of this study given the computational cost. However, based on knowledge of thermal IR (e.g. TES) and total column (e.g. GOSAT) retrievals, use of a single averaging kernel is a reasonable approximation as our study is constrained to Northern Hemisphere summertime where the temperature and sunlight conditions provide sufficient signal for the present evaluation, and because our study looks at the relative merits of different observing approaches.

The prior methane emissions we use are from the EDGARv4.2 anthropogenic methane inventory (European Commission, 2011), the wetland model from Kaplan (2002) as implemented by Pickett-Heaps et al. (2011), the GFED3 biomass burning inventory (van der Werf et al., 2010), a termite inventory and soil absorption from Fung et al. (1991), and a biofuel inventory from Yevich and Logan (2003). Figure 3 shows the total average daily prior methane emissions for the entire North America nested domain. Strong hotspots of CH_4 sources clearly appear over the Canadian wetlands, the Appalachian Mountains (an extensive coal mining area) and densely urbanized areas (e.g., southern California and the East Coast). Following previous assessments of the range of the prior error (Wecht et al., 2014a; Turner et al., 2015). We assume a relative prior standard error of 40% for our bottom-up emission inventory in every grid-cell. This results in a 2.9 Tg/month uncertainty in the total emission budget over North America, a magnitude comparable to the correction to the prior budget found in the inversion of Turner et al. (2015) of 2.3 Tg/month. We assume no prior error correlations, which means that the matrix \mathbf{B} in Eq. 1 is diagonal. Accurately defining error correlations in bottom-up inventories is a challenging problem due to the sparsity of available flux measurements, and is beyond the scope of our study. However, it is likely that the diagonal \mathbf{B} assumption made in our study is overly optimistic, resulting in an overestimation of the spatial resolution of the constraints afforded by the satellite measurements.

2.3 Observations and model uncertainties

We consider several instrument configurations for our study, which are associated with different vertical sensitivities. Constraints on CH_4 emissions are evaluated for the following CH_4 retrievals: the Greenhouse gases Observing SATellite (GOSAT) Proxy XCH4 v3.2 data described by Parker et al. (2011) (available from <http://www.leos.le.ac.uk/GHG/data/>), which consists of CH_4 column mixing ratio X_{CH_4} obtained from Short Wave Infrared (SWIR) measurements near 1.6 μm ; the Tropospheric Emission Spectrometer (TES) V005 Lite product (Worden et al., 2012) (<http://tes.jpl.nasa.gov/data/>), which consists of CH_4 vertical profile retrievals from Thermal Infra Red (TIR) measurements at 7.58–8.55 μm ; and a hypothetical multi-spectral CH_4 profile retrieval, which allows to capture sig-

nal in the boundary layer. Since the DOF for the TES retrievals is less than 2, we use a pressure-weighted TES X_{CH_4} column instead of the retrieved CH_4 profiles. A multi-spectral averaging kernel can be derived by first combining the Jacobians (or sensitivities) of the modeled radiances to methane concentrations from the 1.6 micron and 8 micron bands. Both the TES and GOSAT retrievals also simultaneously estimate interferences such as clouds, albedo, emissivity, temperature, and H_2O . The effects of these interferences can be included by further combining their corresponding Jacobians with the methane Jacobians (e.g., Worden et al., 2004; Kulawik et al., 2006; Butz et al., 2010). Constraints for methane and the other radiative interferences are described in Worden et al. (2012) and Parker et al. (2011). The combination of these Jacobians and constraints are then used to calculate the averaging kernel. The methane component of the resulting multi-spectral, multi-species averaging kernel is then used for this study. The effect of the interferences with this simultaneous retrieval approach is to reduce the overall sensitivity to methane but improve the posteriori errors. A proof of concept for combining near IR and IR-based methane estimates to derived a lower tropospheric estimate is discussed in Worden et al. (2015) using GOSAT and TES profile retrievals.

Figure 2 shows the column averaging kernel for the GOSAT and TES X_{CH_4} retrievals as well as the averaging kernels at three different levels for the multi-spectral retrieval. The GOSAT retrieval sensitivity is nearly uniform throughout the troposphere, with averaging kernel values close to 1. The TES retrieval is mostly sensitive to CH_4 concentrations in the upper troposphere, with a peak of the column averaging kernel around 300 hPa. The multi-spectral profile retrieval shows a distinct signal in the boundary layer, with weaker sensitivities above.

Observation and model transport errors are assumed to be independent and therefore added in quadrature to define the error covariance matrix \mathbf{R} in Eq. 1. Observational errors for GOSAT X_{CH_4} columns are uniformly set to 12 ppb, within the range of values reported in Parker et al. (2011). For the TES retrievals, the profile error covariance matrix is averaged vertically using pressure-weighted functions to obtain X_{CH_4} column errors, as described in Connor et al. (2008). This results in 0.5-2% (or 10-40 ppb) uncertainties for the TES columns (Worden et al., 2012). For the multi-spectral retrievals, a vertically resolved error covariance matrix is used. For comparison, the resulting pressure-weighted column X_{CH_4} error is similar to the one obtained for GOSAT retrievals (~ 12 ppb).

As shown by Locatelli et al. (2013), taking into account transport errors is critical in order to mitigate uncertainties in the inversion, since neglecting them can lead to discrepancies in the posterior estimates of more than 150% of the prior flux at model grid scale. We estimate model transport error using model-data comparison statistics for North American *in situ* observations from the NOAA/ESRL surface, tower, and flask network as well as observations from the HIPPO and CalNex measurement campaigns (Turner et al., 2015). Model error standard deviations are set to 46 ppb in the boundary layer, and 22 ppb in the free-troposphere. Vertical error correlations between simulated concentrations are difficult to quantify with the limited observational sampling available *in*

situ. Transport error correlations between the boundary layer and the free-troposphere are assumed
240 to be negligible due to the decoupling of the physical processes between those two regions. However,
within both the boundary layer and the free-troposphere, a model error correlation of one is assumed
between all altitude levels, which is a conservative (pessimistic) assumption. Our gradient-based
estimates of the inverse Hessian matrix involve generating random perturbations that follow the ob-
servational error statistics (see Section 2.1). For the multi-spectral configuration, a Singular Value
245 Decomposition (SVD) is first performed on the vertically resolved matrix \mathbf{R} in order to generate
independent perturbations (e.g., Bousseret et al., 2015).

In order to assess the relative impact of measurement sensitivity versus spatio-temporal sampling
on the CH_4 emission constraints, both LEO and GEO orbit configurations are considered in our
study. The LEO orbit configuration follows GOSAT's sun-synchronous polar orbit with an Equa-
250 tor overpass local time of $\sim 13:00$. Measurements consist of five across-track points separated by
 ~ 100 km, with footprint diameters of 10.5 km. The GEO configuration corresponds to hourly ob-
servations over North America from 10 to 60° N. The GEO footprint considered is ~ 4 km, therefore
much finer than the GEOS-Chem resolution used (~ 50 km). For both LEO and GEO configurations,
observations are therefore averaged together within each GEOS-Chem grid cell and the instrument
255 error is reduced by multiplying it by the square root of the number of observations.

Finally, contamination by clouds is taken into account for each grid-cell by removing a fraction of
the total number of observations within that cell which corresponds to the GEOS-5 cloud fraction.
The resulting spatial distribution of the observational data density for each satellite configuration
(LEO or GEO) is shown in Fig. 3.

260 3 Results

In the following experiments, we consider the inversion of 30 day, 7 day, and 3 day grid-scale emis-
sion scaling factors over North America. In particular, this means that the spatiotemporal variability
of the methane fluxes (e.g., diurnal cycle and spatial distribution) within each time-window is as-
sumed to be known, and only its magnitude is adjusted. The information content of the inversion is
265 analyzed for four different observational systems:

- a GOSAT instrument onboard a low-Earth orbit platform (GOSAT_LEO)
- a GOSAT instrument onboard a geostationary orbit platform (GOSAT_GEO)
- a TES instrument onboard a geostationary orbit platform (TES_GEO)
- a multi-spectral instrument onboard a geostationary orbit platform (MULTI_GEO).

270 3.1 Error reduction of optimized methane emissions

Figures 4, 5, and 6 show the relative error variance reduction in the emission scaling factors for a 30 day, 7 day, and 3 day inversion, respectively, for each of the observational configurations described above. The DOF, which quantifies the number of pieces of information independently constrained by the observations, is also indicated. For the monthly inversion, the GOSAT_LEO, 275 GOSAT_GEO, and MULTI_GEO configurations show error reductions close to 100% for sparse hotspots over the continent, in particular in the Los Angeles basin, the Central US, the Toronto urban area, the Appalachian Mountains, and the Northeast US. The TES_GEO configuration still shows significant observational constraints in those locations, with error reductions $> 70\%$. However, overall the error reductions afforded by using a TES-like instrument in geostationary orbit are 280 much smaller than the one obtained from a GOSAT-like or multi-spectral instrument. In particular, the DOF for the TES_GEO configuration (164) is about half that of the GOSAT_LEO configuration (298). This demonstrates that using measurements with significant sensitivities to lower-tropospheric concentrations is critical to obtain surface flux information, even in a geostationary framework with high-frequency temporal sampling. The advantage of the GEO over the LEO configuration is more 285 pronounced when smaller emission time-scales are constrained (weekly, 3-day). In particular, the DOF for GOSAT_LEO varies from 88% to 43% of the DOF for GOSAT_GEO between the monthly and the 3-day inversion. Similarly, but to a lesser extent, the benefit of a multi-spectral profile observation compared to a GOSAT-like column measurement is most evident when the temporal resolution of the flux inversion is increased, with a DOF ratio between GOSAT_GEO and MULTI_GEO 290 varying from 84% to 67% between the monthly and 3-day inversions.

These results are synthesized in Figure 7, which shows the relative error variance reduction as a function of emission magnitude, for each observational system and inversion time-window. The convergence of the flux constraints provided by the GOSAT (LEO or GEO) and the multi-spectral GEO instruments is well illustrated by the convergence of the corresponding curves as the tem- 295 poral scale of the optimization increases from 3-day to one-month. These results also show that for grid-cells with high CH_4 emissions ($> 4 \times 10^5 \text{ kg day}^{-1} \text{ grid}^{-1}$), a multi-spectral instrument in geostationary orbit would reduce prior flux errors by more than 80% at time scales as small as 3 days. In particular, this could provide valuable information to monitor the variation of CH_4 emission hotspot activities between workweek and weekend. Finally, we note that Turner et al. (2015) 300 obtained a DOF of 39 for a multi-year CH_4 flux inversion over North America using GOSAT LEO observations, which contrasts with the much higher DOF (298) obtained for our monthly inversion. However, in Turner et al. (2015), a prior dimension reduction of the inverse problem was performed to enable an analytical computation of the solution with only 369 control vector elements. Although it is claimed that the aggregation scheme used to define the reduced space is designed to account 305 for prior error correlations, the results obtained in Turner et al. (2015) indicate the reduction method is suboptimal (see interactive discussion of Turner et al. (2015) for more details) and there is no

guarantee that the implicitly defined correlations are realistic. On the other hand, in our case neglecting error correlations in the prior inventory results in an overestimation of the DOF. In the absence of a rigorous methodology to accurately estimate the prior error correlations, the DOFs we derived should be interpreted as an upper-limit for the different observational configurations, to which one can tend as the spatial and temporal resolutions of the bottom-up CH₄ inventories are refined. In relation to previous works by Turner et al. (2015) and Bocquet (2011), it should also be noted that the gradient-based algorithm used in our study allows us to estimate the DOF of the inversion prior to optimization; this information could therefore be used to objectively determine an appropriate dimension for the inverse problem, upon which specific dimension reduction methods could be devised.

3.2 Spatial resolution of the inversion

An objective measure of the spatial resolution of the inversion, i.e., the ability of the observational system to constrain grid-scale emissions independently from each other, is provided by the rows of the model resolution matrix (see Eq. 5). Figure 8 shows the model resolution matrix rows of the weekly inversion corresponding to five different locations, chosen to span a range of characteristics, in terms of emissions magnitude and error reduction. For readability, only grid-cells included within the largest circle centered on each location and containing values greater than 0.05 are shown. Table 1 summarizes the coordinates and CH₄ emissions corresponding to each location. Since the model grid-cell area depends on the latitude, the radiuses of each of the structures shown in Figure 8 are also summarized in Table 2. Note that the 3-day inversion results (not shown) gave similar results as the one-week inversion. The gain in spatial resolution of the optimized fluxes when a GEO orbit is used is evident when comparing the GOSAT_LEO and GOSAT_GEO results. In particular, Table 2 suggests that for the Central US and California regions, the spatial resolution of the independently constrained flux patterns is about two times higher in the case of a GEO configuration (radius~80 km) compared to a LEO configuration (radius~160 km). Based on the comparison between the GOSAT_GEO and MULTI_GEO configurations, the gain in spatial resolution afforded by the use of a multi-spectral instrument appears significant (factor of 2) only over the Eastern US region. Note that although the size of the flux structures are similar between the TES_GEO and GOSAT_LEO configurations, the average values of the model resolution matrix row within each structure are significantly higher in the case of GOSAT_LEO.

3.3 Impact of boundary and initial conditions uncertainties

Boundary and initial conditions used in the forward transport model contain errors. Therefore, any consistent flux inversion system should jointly optimize the fluxes, initial state and boundary conditions. However, in practice many studies overlook this issue and optimize those quantities separately (e.g., Basu et al., 2013; Wecht et al., 2014a; Deng et al., 2014). In the latter case, a flux-only inver-

sion is performed with initial and boundary conditions that are effectively assumed perfectly known. It is therefore of interest to estimate the impact of errors in the initial and boundary conditions on the optimized fluxes. Figure 9 shows the perturbations in the optimized emission scaling factors for the weekly inversion resulting from random gaussian perturbations of the boundary conditions with standard deviation 16 ppb. The choice for the standard error of the noise is based on model-data comparisons from the HIAPER Pole-to-Pole Observations (HIPPO) experiment (Turner et al., 2015), which consists in extensive aircraft measurements throughout the troposphere over the Pacific ocean. Only weekly inversion results are shown here so that enough constraints are obtained for all observational configurations while keeping the computational cost the inversions manageable.

For all configurations, the results show scaling factor perturbations throughout the North America domain, although they are less pronounced over the Eastern US due to the dominant westerly propagation of the boundary condition perturbations into the domain. The TES_GEO and GOSAT_GEO configurations show similar sensitivities of the optimized scaling factors to boundary conditions, with large areas characterized by perturbations between 10% and 50%, and with impacts greater than 50% locally. In comparison, the GOSAT_GEO configuration shows smaller sensitivities to boundary conditions, with perturbations generally smaller than 30%. The MULTI_GEO results are in contrast with the other configurations, with most scaling factor perturbations being smaller than 5%.

The differences between the sensitivities of the optimized fluxes to boundary conditions for different observational systems are driven by two factors: 1) the sensitivity of the observations to the underlying fluxes (defined by the operator H) and 2) the model-data mismatch (i.e., $H(\mathbf{x}) - \mathbf{y}$). This can be seen, e.g., by considering the observational term in the gradient formula of Eq. 2. Formally, a perturbation of the boundary conditions will translate into a corresponding perturbation of the observations (\mathbf{y}) in the model-data mismatch, which is propagated into flux scaling factor perturbations through the adjoint matrix of sensitivities (\mathbf{H}^T). The effect of 1) is clearly seen when comparing the GOSAT_GEO and GOSAT_LEO results, the higher temporal frequency of the geostationary observations providing higher sensitivity to the fluxes. The effect of 2) is best illustrated by comparing the GOSAT_GEO and MULTI_GEO results. Indeed, since the multi-spectral measurements allow for distinguishing boundary layer from free tropospheric CH_4 concentrations, and given the uniform (~ 1) sensitivity of the GOSAT column measurements throughout the troposphere (see Figure 2), the boundary layer model-data mismatch (MULTI_GEO) is much smaller than the column model-data mismatch (GOSAT_GEO), which results in much higher flux adjustments for the GOSAT_GEO configuration.

The same analysis applies to the sensitivities of the optimized fluxes to initial conditions, which are shown in Figure 10. Here the CH_4 3D initial concentrations were perturbed with random gaussian noises of standard deviation 46 ppb and 22 ppb in the boundary layer and the free troposphere, respectively, based on model-data comparisons with NOAA flasks, tall tower, and aircraft measurements over North America (Turner et al., 2015). In the case of initial conditions, as opposed to

boundary conditions, the forcing perturbations are applied only once at the beginning of the inversion window, which results in the signal being quickly “diluted” and therefore in smaller impacts on the optimized fluxes. The GOSAT_GEO configuration, which combines significant sensitivities to CH₄ concentrations throughout the troposphere with high-frequency measurements, is most sensitive to initial condition perturbations, with up to 30% variability in the optimized scaling factors. The GOSAT_LEO and TES_GEO configurations show comparable sensitivities, with scaling factors perturbations generally smaller than 10%. Similarly to the boundary condition case, initial condition sensitivities associated with the MULTI_GEO configuration are about one order of magnitude smaller than other configurations, with scaling factors perturbations generally smaller than 3%. These results show that although the advantage of a multi-spectral instrument in term of spatiotemporal constraints on the fluxes becomes significant only for time-scales smaller than a week, there is still a clear benefit in using this configuration to mitigate the impact of uncertainties in boundary and initial conditions on the inversion, even when optimizing fluxes at coarser temporal resolution (e.g., weekly or monthly).

4 Conclusions

In this paper we evaluated top-down constraints on methane emissions in North America provided by potential future geostationary observation missions (GEO-CAPE) and existing low-earth orbit remote sensing instruments (GOSAT). For the first time, a grid-scale estimate of the information content of a high-resolution inversion ($0.5^\circ \times 0.7^\circ$ over North America) in a 4D-Var inversion framework has been performed using an efficient stochastic algorithm. In particular, this allowed us to compute both the relative error reductions and the spatial correlations between observational constraints in the inversion. Instrument configurations corresponding to current TIR and SWIR methane products (TES and GOSAT, respectively), as well as a potential multi-spectral retrieval, were considered. This allowed us to assess the relative importance of the vertical sensitivity of the measurement versus the spatiotemporal resolution of the sampling (GEO versus LEO) in methane flux inversions.

We found that a GEO configuration provides significant benefits over current LEO products in term of error reductions in the optimized fluxes when the targeted time-scales are about a week or less. For a 3-day inversion, the number of pieces of information (DOF) independently constrained by the GEO observations is about twice as many than in the case of a LEO configuration (DOF of 266 and 115, respectively). Experiments with TIR GEO and SWIR LEO configurations demonstrated that the high temporal frequency of GEO observations cannot compensate for weak sensitivities of the satellite measurement to boundary layer concentrations, since constraints from a TES-like instrument on GEO orbit correspond to only about a half of the information content afforded by a GOSAT instrument on LEO orbit for a monthly inversion (DOF of 164 and 298, respectively). The benefit of using a multi-spectral instrument in flux inversions compared to a SWIR instrument

has been demonstrated for weekly to sub-weekly scale constraints, with an increase in the DOF of
415 about 50% for a 3-day inversion. For the multi-spectral GEO configuration, the information content
is similar for a 3-day or a one-month optimization (DOFs of 397 and 398, respectively).

Over some local CH₄ source hotspots (emissions > 4 × 10⁵ kg day⁻¹) in the Central US, Cal-
ifornia and Eastern US, both SWIR and multi-spectral GEO configurations allow for nearly com-
plete constraints on emissions (error reduction close to 100%) at spatial resolution smaller than
420 100km×100km. These estimates are optimistic, given the lack of spatial error correlation consid-
ered in our prior emissions, which should be addressed in future work, but do reveal the potential
spatial resolution provided by the measurements alone.

The sensitivity of the optimized emission scaling factors to uncertainties in initial and boundary
conditions has also been assessed by propagating random perturbations of these forcings into the flux
425 estimates. While the flux responses to the boundary and initial condition perturbations can reach 50%
and 30%, respectively, in the case of GOSAT column constraints, they were an order of magnitude
lower (<5%) in the case of multi-spectral profile observations.

With growing concerns about the environmental impacts of CH₄ emissions from the oil and gas
industry and the urge for better monitoring of the nation's CH₄ budget, a multi-spectral instru-
430 ment onboard geostationary orbit would provide a key tool to characterize CH₄ fluxes' variability at
weekly to sub-weekly time-scale, while greatly mitigating the impact of inverse method uncertainties
on the optimized fluxes. Moreover, such an observational system would allow for better understand-
ing of the critical role of wetlands in the global methane budget and their impact on climate change
(e.g., Bloom et al., 2012; Miller et al., 2014). Further investigations would be needed to quantify
435 the sensitivity of these results to the choice of the reference CH₄ emission inventory, since signifi-
cant discrepancies in the magnitude and spatiotemporal distributions of CH₄ sources exist between
current bottom-up inventory (Kirschke et al., 2013).

In our study we have neglected prior error correlations in the absence of robust data and methodol-
ogy to rigorously estimate them. Since error correlations in prior bottom-up inventories nevertheless
440 exist, additional experiments should be performed to test the sensitivity of our information con-
tent analysis to different error correlation structures. Likewise, spatial correlations associated with
model and observation errors should be included in future OSSEs in order to obtain more reliable er-
ror reduction estimates. We have also performed the inversion using emission scaling factors, which
effectively places a hard constraint on the spatial distribution of the emissions - an assumption that
445 warrants further investigations. The robustness of our results against model and observational biases
should also be investigated. Finally, following recent studies investigating regional to urban con-
straints from geostationary remote-sensing instruments (Polonsky et al., 2014; Rayner et al., 2014),
it would be interesting to apply the present methodological framework to inversions at much higher
spatiotemporal resolution in order to analyze the ability of such observational system to extract in-
450 formation at spatial scales of only a few km².

Acknowledgements. This project was supported by the NASA GEO-CAPE Science Team grant NNX14AH02G and the NOAA grant NA14OAR4310136. This work utilized the Janus supercomputer, which is supported by the National Science Foundation (award number CNS-0821 794) and the University of Colorado Boulder. The Janus supercomputer is a joint effort of the University of Colorado Boulder, the University of Colorado Denver and the National Center for Atmospheric Research. Alexander J. Turner was supported by a Department of Energy (DOE) Computational Science Graduate Fellowship (CSGF). Part of this research was carried out at the Jet Propulsion Laboratory, California Institute of Technology, under a contract with the National Aeronautics and Space Administration.

References

- 460 Basu, S., Guerlet, S., Butz, A., Houweling, S., Hasekamp, O., Aben, I., Krummel, P., Steele, P., Langenfelds, R.,
Torn, M., Biraud, S., Stephens, B., Andrews, A., and Worthy, D.: Global CO₂ fluxes estimated from GOSAT
retrievals of total column CO₂, *Atmos. Chem. Phys.*, 13, 8695–8717, doi:10.5194/acp-13-8695-2013, 2013.
- Bloom, A. A., Palmer, P. I., Fraser, A., and Reay, D. S.: Seasonal variability of tropical wetland CH₄ emissions:
the role of the methanogen-available carbon pool, *Biogeosciences*, 9, 2821–2830, doi:10.5194/bg-9-2821-
465 2012, 2012.
- Bousserez, N., Henze, D. K., Perkins, A., Bowman, K. W., Lee, M., Liu, J., Deng, F., and Jones, D. B. A.:
Improved analysis-error covariance matrix for high-dimensional variational inversions: application to source
estimation using a 3-D atmospheric transport model, *Q. J. Roy. Meteor. Soc.*, doi:10.1002/qj.2495, 2015.
- Butz, A., Hasekamp, O. P., Frankenberg, C., Vidot, J., and Aben, I.: CH₄ retrievals from space-based solar
470 backscatter measurements: Performance evaluation against simulated aerosol and cirrus loaded scenes, *J.*
Geophys. Res., 115, D24302, doi:10.1029/2010JD014514.
- Caulton, D. R., Shepson, P. B., Santoro, R. L., Sparks, J. P., Howarth, R. W., Ingraffea, A. R., Cambaliza, M. O.,
Sweeney, C., Karion, A., Davis, K. J., Stirm, B. H., Montzka, S. A., and Miller, B. R.: Toward a better
understanding and quantification of methane emissions from shale gas development, *P. Natl. Acad. Sci.*
475 USA, 111, 6237–6242, 2014.
- Chance, K., Liu, X., Suleiman, R. M., Flittner, D. E., Al-Saadi, J., and Janz, S. J.: Tropospheric emissions:
monitoring of pollution (TEMPO), *P. Soc Photo-Opt. Ins.*, 8866, 88660D, 2013.
- Chevallier, F., Bréon, F. M., and Rayner, P. J.: Contribution of the Orbiting Carbon Observatory to the estimation
of CO₂ sources and sinks: Theoretical study in a variational data assimilation framework, *J. Geophys. Res.-*
480 *Atmos.*, 112(D9),2156–2202, 2007.
- Chevallier, F., Wang, T., Ciais, P., Maignan, F., Bocquet, M., Altaf Arain, M., Cescatti, A., Chen, J., Dolman, A.,
Beverly, E., Margolis, H. A., Montagani, L., and Moors, .E. J.: What eddy-covariance measurements tell us
about prior land flux errors in CO₂-flux inversion schemes, *Global Biogeochemical Cycles*, 26, 1,1994-9224,
2012.
- 485 Connor, B. J., Boesch, H., Toon, G., Sen, B., Miller, C., and Crisp, D.: Orbiting Carbon Observatory: Inverse
method and prospective error analysis, *J. Geophys. Res.-Atmos.*, 113, D05305, doi:10.1029/2006JD008336,
2008.
- Cressot, C., Chevallier, F., Bousquet, P., Crevoisier, C., Dlugokencky, E. J., A., Fortems-Cheiney, A., Franken-
berg, C., Parker, R., Pison, I., Scheepmaker, R. A., Montzka, S. A., Montzka, S. A., Krummel, P. B.,
490 Steele, L. P., and Langenfelds, R. L.: On the consistency between global and regional methane emissions
inferred from SCIAMACHY, TANSO-FTS, IASI and surface measurements, *Atmos. Chem. Phys.*, 14, 577–
592, doi:10.5194/acp-14-577-2014, 2014.
- Deng, F., Jones, D. B. A., Henze, D. K., Bousserez, N., Bowman, K. W., Fisher, J. B., Nassar, R., O’Dell, C.,
Wunch, D., Wennberg, P. O., Kort, E. A., Wofsy, S. C., Blumenstock, T., Deutscher, N. M., Griffith, D. W. T.,
495 Hase, F., Heikkinen, P., Sherlock, V., Strong, K., Sussmann, R., and Warneke, T.: Inferring regional
sources and sinks of atmospheric CO₂ from GOSAT XCO₂ data, *Atmos. Chem. Phys.*, 14, 3703-3727,
doi:10.5194/acp-14-3703-2014, 2014.

- Dlugokencky, E. J., Nisbet, E. G., Fisher, R., and Lowry, D.: Global atmospheric methane: budget, changes and dangers, *Philos. T. R. Soc. A*, 369, 2058–2072, 2011.
- 500 Fiore, A. M., Jacob, D. J., Field, B. D., Streets, D. G., Fernandes, S. D., and Jang, C.: Linking ozone pollution and climate change: the case for controlling methane, *Geophys. Res. Lett.*, 29(19), 25–1, 2002.
- Fiore, A. M., West, J. J., Horowitz, L. W., Naik, V., and Schwarzkopf, M. D.: Characterizing the tropospheric ozone response to methane emission controls and the benefits to climate and air quality, *J. Geophys. Res.-Atmos.*, 113(D8),1984–2012, 2008.
- 505 Fishman, J., Iraci, L., Al-Saadi, J., Chance, K., Chavez, F., Chin, M., Coble, P., Davis, C., DiGiacomo, P., Edwards, D., Eldering, L., Goes, J., Herman, J., Hu, C., Jacob, D. J., Jordan, C., Kawa, S. R., Key, R., Liu, X., Lohrenz, S., Mannino, A., Natraj, V., Neil, D., Neu, J., Newchurch, M., Pickering, K., Salisbury, J., Sosik, H., Subramaniam, A., Tzortziou, M., Wang, J. and Wang, M.: The United States’ next generation of atmospheric composition and coastal ecosystem measurements: NASA’s Geostationary Coastal and Air
- 510 Pollution Events (GEO-CAPE) Mission, *B. Am. Meteorol. Soc.*, 93, 1547–1566, 2012.
- Fung, I., John, J., Lerner, J., Matthews, E., Prather, M., Steele, L., and Fraser, P.: Three-dimensional model synthesis of the global methane cycle, *J. Geophys. Res.-Atmos.*, 96, 13033–13065, 1991.
- Gazovic, M., Kutzbach, L., Schreiber, P., Wille, C., and Wilmking, M.: Diurnal dynamics of CH₄ from a boreal peatland during snowmelt, *Tellus B*, 62, 133–139, 2010.
- 515 Kaplan, J. O.: Wetlands at the Last Glacial Maximum: distribution and methane emissions, *Geophys. Res. Lett.*, 29, doi: 10.1029/2001GL013366, 2002.
- Karion, A., Sweeney, C., Pétron, G., Frost, G., Michael Hardesty, R., Kofler, J., Miller, B. R., Newberger, T., Wolter, S., Banta, R., Brewer, A., Dlugokencky, E., Lang, P., Montzka, S. A., Schnell, R., Tans, P., Trainer, M., Zamora, R. and Conley, S: Methane emissions estimate from airborne measurements over a western United States natural gas field, *Geophys. Res. Lett.*, 40, 4393–4397, 2013.
- 520 Katzenstein, A. S., Doezema, L. A., Simpson, I. J., Blake, D. R., and Rowland, F. S.: Extensive regional atmospheric hydrocarbon pollution in the southwestern United States, *P. Natl. Acad. Sci. USA*, 100, 11975–11979, 2003.
- Kirschke, S., Bousquet, P., Ciais, P., Saunois, M., Canadell, J. G., Dlugokencky, E. J., Bergamaschi, P., Bergmann, D., Blake, D. P., Bruhwiler, L., Cameron-Smith, P., Castaldi, P., Chevallier, F., Feng, L., Fraser, A., Heimann, M., Hodson, E. L., Houweling, S., Josse, B., Fraser, P. J., Krummel, P. B., Lamarque, J. F., Langenfelds, R. L., Le Quere, C., Naik, V., O’Doherty, S., Palmer, P. I., Pison, I., Plummer, D., Poulter, B., Prinn, R. G., Rigby, M., Ringeval, B., Santini, M., Schmidt, M., Shindell, D. T., Simpson, I. J., Spahni, R., Steele, L. P., Strode, S. A., Sudo, K., Szopa, S., van der Werf, G. R., Voulgarakis, A., van
- 530 Weele, M., Weiss, R. F., Williams, J. E., Zen, G.: Three decades of global methane sources and sinks, *Nat. Geosci.*, 6, 813–823, 2013.
- Kort, E. A., Eluszkiewicz, J., Stephens, B. B., Miller, J. B., Gerbig, C., Nehr Korn, T., Daube, B. C., Kaplan, J. O., Houweling, S., and Wofsy, S. C.: Emissions of CH₄ and N₂O over the United States and Canada based on a receptor-oriented modeling framework and COBRA-NA atmospheric observations, *Geophys. Res. Lett.*, 35, 2008.
- 535 Kulawik, S. S., Worden, J., Eldering, A., Bowman, K., Gunson, M., Osterman, G. B., Zhang, L., Clough, S., Shephard, M. W. and Beer, R.: Implementation of cloud retrievals for Tropospheric Emission Spectrometer

- (TES) atmospheric retrievals: part 1. Description and characterization of errors on trace gas retrievals, *J. Geophys. Res.*, 111, D24204, doi:10.1029/2005JD006733.
- 540 Lions, J. L.: *Optimal Control of Systems Governed by Partial Differential Equations*, Springer-Verlag, Berlin, 1971.
- Ben R. Milleri, John B. Milleri, Stephen A. Montzka, Thomas Nehrkorn, and Colm Sweeney
- Liu, J., Bowman, K., Lee, M., Henze, D. K., Bousseres, N., Brix, H., Collatz, G., Menemenlis, D., Ott, L., Pawson, S., Jones, D., and Nassar, R.: Carbon monitoring system flux estimation and attribution: impact of
- 545 ACOS-GOSAT XCO₂ sampling on the inference of terrestrial biospheric sources and sinks, *Tellus B*, 62, 0, 2014.
- Locatelli, R., Bousquet, P., Chevallier, F., Fortems-Cheney, A., Szopa, S., Saunois, M., Agusti-Panareda, A., Bergmann, D., Bian, H., Cameron-Smith, P., Chipperfield, M. P., Gloor, E., Houweling, S., Kawa, S. R., Krol, M., Patra, P. K., Prinn, R. G., Rigby, M., Saito, R., and Wilson, C.: Impact of transport model errors
- 550 on the global and regional methane emissions estimated by inverse modelling, *Atmos. Chem. Phys.*, 13, 9917–9937, doi:10.5194/acp-13-9917-2013, 2013.
- Meirink, J. F., Bergamaschi, P., and Krol, M. C.: Four-dimensional variational data assimilation for inverse modelling of atmospheric methane emissions: method and comparison with synthesis inversion, *Atmos. Chem. Phys.*, 8, 6341–6353, doi:10.5194/acp-14-6341-2008, 2008.
- 555 Miller, S. M., Wofsy, S. C., Michalak, A. M., Kort, E. A., Andrews, A. E., Biraud, S. C., Dlugokencky, E. J., Eluszkiewicz, J., Fischer, M. L., Janssens-Maenhout, G., Miller, B. R., Miller, J. B., Montzka, S. A., Nehrkorn, T., and Sweeney, C.: Anthropogenic emissions of methane in the United States, *P. Natl. Acad. Sci. USA*, 110, 20018–20022, 2013.
- Miller, S. M., Worthy, D. E., Michalak, A. M., Wofsy, S. C., Kort, E. A., Havice, T. C., Andrews, A. E.,
- 560 Dlugokencky, E. J., Kaplan, J. O., Levi, P. J., Tian, H., and Zhang, B.: Observational constraints on the distribution, seasonality, and environmental predictors of North American boreal methane emissions, *Global Biogeochem. Cy.*, 28, 146–160, 2014.
- Morin, T. H., Bohrer, G., Naor-Azrieli, L., Mesi, S., Kenny, W. T., Mitsch, W. J., and Schaefer, K. V. R.: The seasonal and diurnal dynamics of methane flux at a created urban wetland, *Ecol. Eng.*, 72, 74–83, 2014.
- 565 Myhre, G. and Shindell, D.: *Climate Change 2013: The Physical Science Basis*, Intergovernmental Panel on Climate Change (IPCC), Chap. 8, 2013.
- Parker, R., Boesch, H., Cogan, A., Fraser, A., Feng, L., Palmer, P. I., Messerschmidt, J., Deutscher, N., Griffith, D. W., Notholt, J., Wennberg, P. O., and Wunch, D.: Methane observations from the Greenhouse Gases Observing SATellite: comparison to ground-based TCCON data and model calculations, *Geophys. Res. Lett.*, 38, L15807, doi:10.1029/2011GL047871, 2011.
- 570 Pickett-Heaps, C. A., Jacob, D. J., Wecht, K. J., Kort, E. A., Wofsy, S. C., Diskin, G. S., Worthy, D. E. J., Kaplan, J. O., Bey, I., and Drevet, J.: Magnitude and seasonality of wetland methane emissions from the Hudson Bay Lowlands (Canada), *Atmos. Chem. Phys.*, 11, 3773–3779, doi:10.5194/acp-11-3773-2011, 2011.
- Polonsky, I. N., O'Brien, D. M., Kumer, J. B., O'Dell, C. W., and the geoCARB Team: sPerformance of a geostationary mission, geoCARB, to measure CO₂, CH₄ and CO column-averaged concentrations, *Atmos. Meas. Tech.*, 7, 959–981, doi:10.5194/amt-7-959-2014, 2014.

- Rayner, P. J., Utembe, S. R., and Crowell, S.: Constraining regional greenhouse gas emissions using geostationary concentration measurements: a theoretical study, *Atmos. Meas. Tech.*, 7, 3285–3293, doi:10.5194/amt-7-3285-2014, 2014.
- 580 Shindell, D., Kuylensstierna, J. C., Vignati, E., van Dingenen, R., Amann, M., Klimont, Z., Anenberg, S. C., Muller, N., Janssens-Maenhout, G., Raes, F., Schwartz, J., Faluvegi, G., Pozzoli, L., Kupiainen, K., Hoglund-Isaksson, L., Emberson, L., Streets, D., Ramanathan, V., Hicks, K., Oanh, N., Milly, G., Williams, M., Demkine, V., and Fowler, D.: Simultaneously mitigating near-term climate change and improving human health and food security, *Science*, 335, 183–189, 2012.
- 585 Tarantola, A.: Inverse problem theory and methods for model parameter estimation, SIAM, <http://dx.doi.org/10.1137/1.9780898717921>, 2005.
- Turner, A. J. and Jacob, D. J.: Balancing aggregation and smoothing errors in inverse models, *Atmos. Chem. Phys. Discuss.*, 15, 1001–1026, doi:10.5194/acpd-15-1001-2015, 2015.
- Turner, A. J., Jacob, D. J., Wecht, K. J., Maasakkers, J. D., Biraud, S. C., Boesch, H., Bowman, K. W.,
590 Deutscher, N. M., Dubey, M. K., Griffith, D. W. T., Hase, F., Kuze, A., Notholt, J., Ohyama, H., Parker, R., Payne, V. H., Sussmann, R., Velazco, V. A., Warneke, T., Wennberg, P. O., and Wunch, D.: Estimating global and North American methane emissions with high spatial resolution using GOSAT satellite data, *Atmos. Chem. Phys. Discuss.*, 15, 4495–4536, doi:10.5194/acpd-15-4495-2015, 2015.
- van der Werf, G. R., Randerson, J. T., Giglio, L., Collatz, G. J., Mu, M., Kasibhatla, P. S., Morton, D. C.,
595 DeFries, R. S., Jin, Y., and van Leeuwen, T. T.: Global fire emissions and the contribution of deforestation, savanna, forest, agricultural, and peat fires (1997–2009), *Atmos. Chem. Phys.*, 10, 11707–11735, doi:10.5194/acp-10-11707-2010, 2010.
- Wecht, K. J., Jacob, D. J., Wofsy, S. C., Kort, E. A., Worden, J. R., Kulawik, S. S., Henze, D. K., Kopacz, M., and Payne, V. H.: Validation of TES methane with HIPPO aircraft observations: implications for inverse
600 modeling of methane sources, *Atmos. Chem. Phys.*, 12, 1823–1832, doi:10.5194/acp-12-1823-2012, 2012.
- Wecht, K. J., Jacob, D. J., Frankenberg, C., Jiang, Z., and Blake, D. R.: Mapping of North American methane emissions with high spatial resolution by inversion of SCIAMACHY satellite data, *J. Geophys. Res.-Atmos.*, 119, 7741–7756, 2014a.
- Wecht, K. J., Jacob, D. J., Sulprizio, M. P., Santoni, G. W., Wofsy, S. C., Parker, R., Bösch, H., and Worden, J.:
605 Spatially resolving methane emissions in California: constraints from the CalNex aircraft campaign and from present (GOSAT, TES) and future (TROPOMI, geostationary) satellite observations, *Atmos. Chem. Phys.*, 14, 8173–8184, doi:10.5194/acp-14-8173-2014, 2014b.
- West, J. J. and Fiore, A. M.: Management of tropospheric ozone by reducing methane emissions, *Environ. Sci. Technol.*, 39, 4685–4691, 2005.
- 610 West, J. J., Fiore, A. M., Horowitz, L. W., and Mauzerall, D. L.: Global health benefits of mitigating ozone pollution with methane emission controls, *P. Natl. Acad. Sci. USA*, 103, 3988–3993, 2006.
- West, J. J., Fiore, A. M., and Horowitz, L. W.: Scenarios of methane emission reductions to 2030: abatement costs and co-benefits to ozone air quality and human mortality, *Climatic Change*, 114, 441–461, 2012.
- Worden, J., Kulawik, S., Shephard, M. W., Clough, S. A., Worden, H., Bowman, K., and Goldman, A.: Predicted
615 errors of tropospheric emission spectrometer nadir retrievals from spectral window selection, *J. Geophys. Res.*, 109, D09308, doi:10.1029/2004JD004522.

- Worden, J., Kulawik, S., Frankenberg, C., Payne, V., Bowman, K., Cady-Peirara, K., Wecht, K., Lee, J.-E., and Noone, D.: Profiles of CH₄, HDO, H₂O, and N₂O with improved lower tropospheric vertical resolution from Aura TES radiances, *Atmos. Meas. Tech.*, 5, 397–411, doi:10.5194/amt-5-397-2012, 2012.
- 620 Worden, J. R., Turner, A. J., Bloom, A., Kulawik, S. S., Liu, J., Lee, M., Weidner, R., Bowman, K., Frankenberg, C., Parker, R., Payne, V. H.: Quantifying lower tropospheric methane concentrations using GOSAT near-IR and TES thermal IR measurements, *Atmos. Meas. Tech.*, 8, 3433–3445, doi:10.5194/amt-8-3433-2015, 2015.
- Xiao, Y., Logan, J. A., Jacob, D. J., Hudman, R. C., Yantosca, R., and Blake, D. R.: Global budget of ethane and regional constraints on US sources, *J. Geophys. Res.-Atmos.*, 113, D21, doi:10.1029/2007JD009415, 2008.
- 625 Yevich, R. and Logan, J. A.: An assessment of biofuel use and burning of agricultural waste in the developing world, *Global Biogeochem. Cy.*, 17, 4, doi:10.1029/2002GB001952, 2003.
- Zavala-Araiza, D. et al: Reconciling divergent estimates of oil and gas methane emissions, *Proc. Natl. Acad. Sci.*, 201522126, doi:10.1073/pnas.1522126112, 2015.
- 630 Zhen, L., Pinto, J. P., Turner, A. J., Henze, D. K., Brioude, J., Bousserez, N., Bruhwiler, L. M. P., and Michelsen, H. P.: Mapping methane emissions from oil and natural gas systems in the contiguous United States, *Environmental Science & Technology*, submitted.
- Zhu, L., Henze, D., Bash, J. O., Cady-Pereira, K. E., Shephard, M. W., Luo, M., and Capps, S. L.: Sources and impacts of atmospheric NH₃: Current understanding and frontiers for modeling, measurements, and remote sensing in North America, *Current Pollution Reports*, in press.
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Table 1. Coordinates of the five locations considered for the rows of the model resolution matrix, with their corresponding emission rate.

Region	Coordinates (lon, lat (°))	Emission ($10^5 \text{ kg day}^{-1} (\text{grid cell})^{-1}$)	Emission ($10^5 \text{ kg day}^{-1} \text{ km}^{-2}$)
Eastern US	(−82, 38)	399	0.12
Central US	(−104, 40)	830	0.26
California	(−117.3, 34.5)	895	0.26
Western Canadian	(−120, 61.5)	575	0.29
Wetlands			
Eastern Canadian	(−84.6, 52.5)	205	0.08
Wetlands			

Table 2. Coordinates of the five locations considered for the rows of the model resolution matrix and approximate radius of influence of neighboring grid cells (see text), for each satellite configuration and a weekly methane flux inversion.

Region	Coordinates (lon, lat (°))	TES_GEO Radius (km)	GOSAT_LEO Radius (km)	GOSAT_GEO Radius (km)	MULTI_GEO Radius (km)
Eastern US	(−82, 38)	160	160	160	80
Central US	(−104, 40)	79	158	79	79
California	(−117.3, 34.5)	164	164	82	82
Western Canadian	(−120, 61.5)	130	196	131	196
Wetlands					
Eastern Canadian	(−84.6, 52.5)	283	213	142	142
Wetlands					

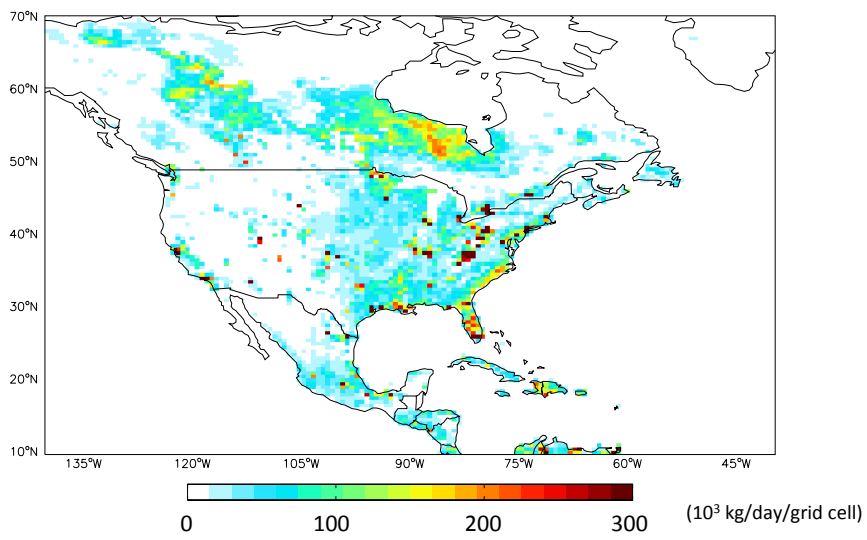


Figure 1. Total daily average prior methane emissions for the nested North America domain ($0.5^\circ \times 0.7^\circ$).

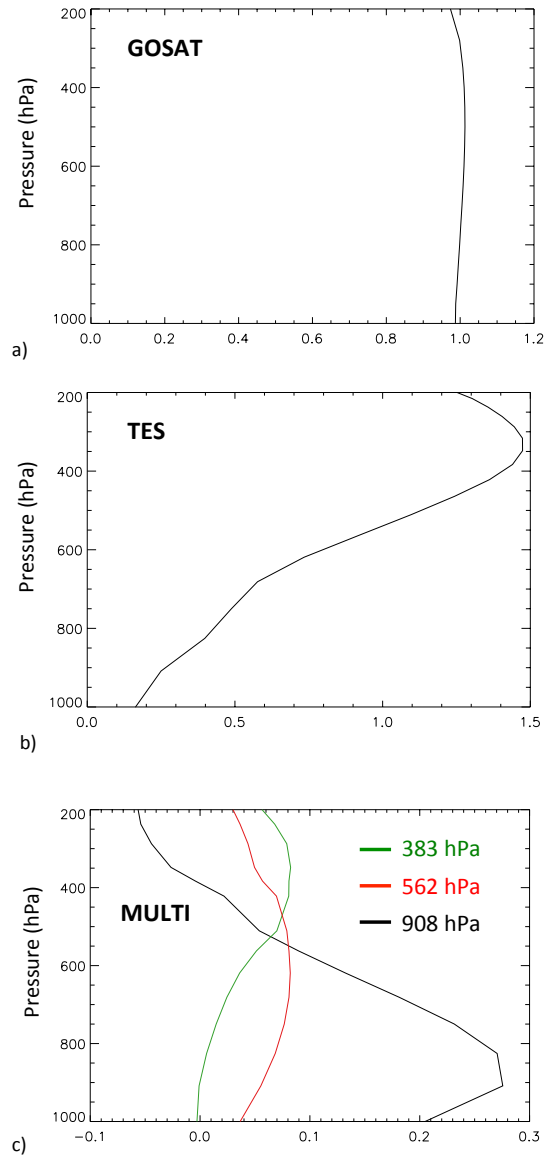


Figure 2. Averaging kernels for the different instrument configurations: **(a)** GOSAT column averaging kernel; **(b)** TES column averaging kernel; **(c)** Multi-spectral averaging kernels at three pressure levels: 908, 562 and 383 hPa.

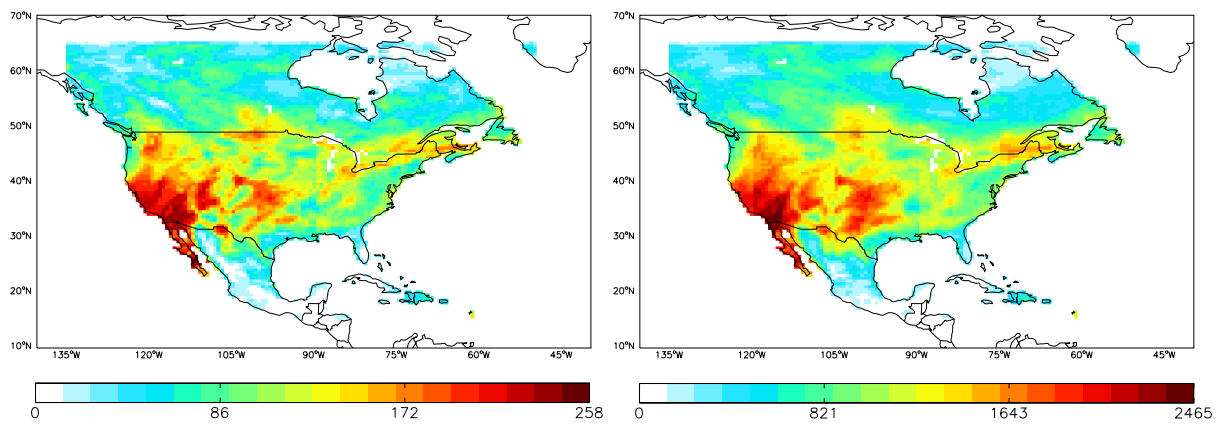


Figure 3. Density of satellite observations (/grid cell/week) for LEO (left) and GEO (right) orbits for the nested North America domain ($0.5^\circ \times 0.7^\circ$) and for the period 1–8 July 2008.

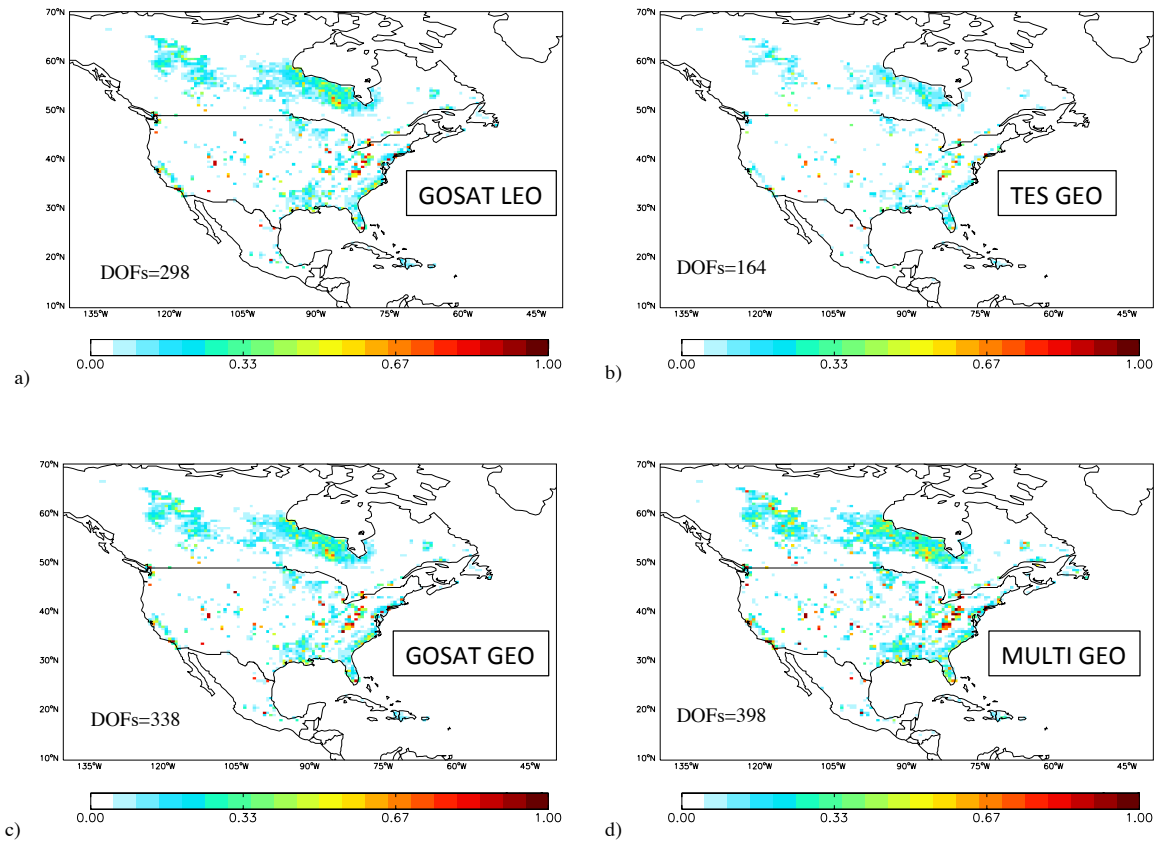


Figure 4. Relative error variance reduction for a 30 day methane emission optimization (1–30 July 2008) using: (a) GOSAT low-earth orbit observations (GOSAT_LEO); (b) GEO-CAPE observations with a TES-like instrument (TES_GEO); (c) GEO-CAPE observations with a GOSAT-like instrument (GOSAT_GEO); (d) GEO-CAPE observations with a multi-spectral instrument (MULTI_GEO). Zero values correspond to emissions with no constraints from observations, while values of one correspond to emissions entirely constrained by observations. The DOF for each inversion, which is the sum of all diagonal elements of the model resolution matrix, is also indicated.

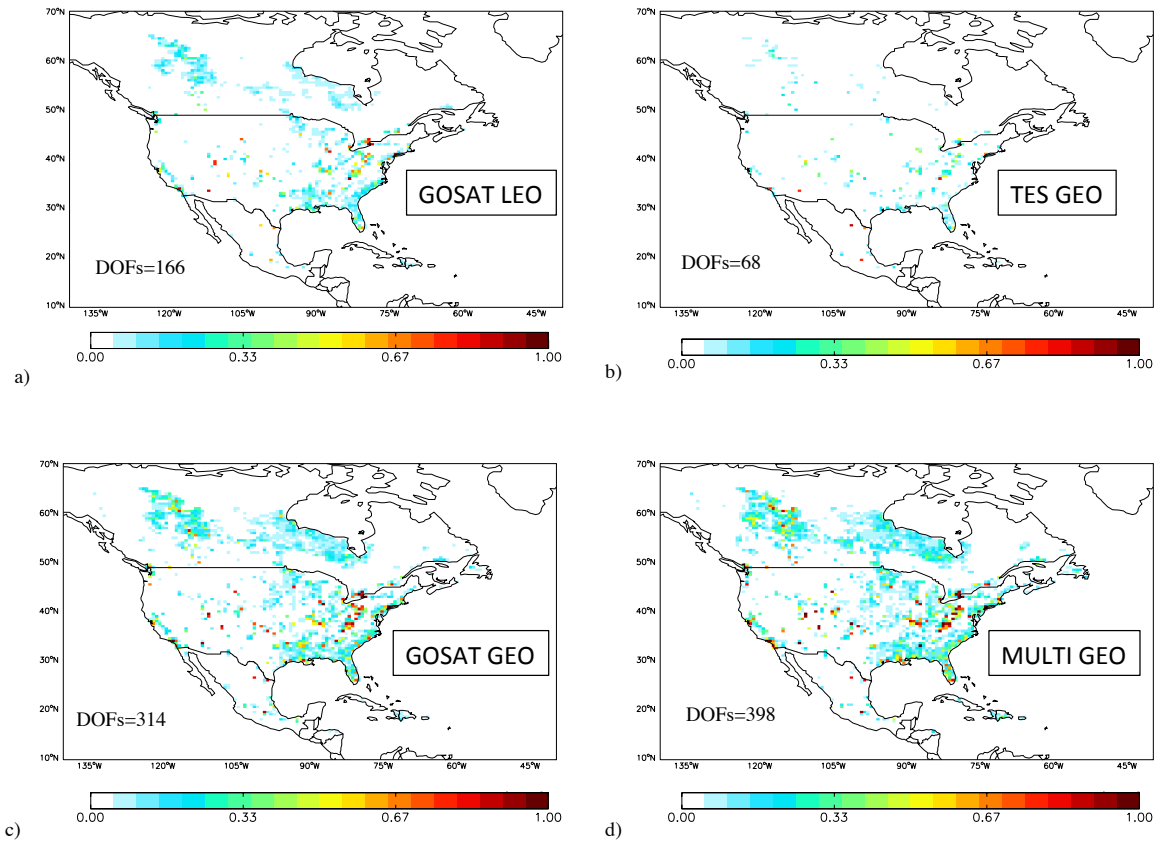


Figure 5. Relative error variance reduction for a 7 day methane emission optimization (1–8 July 2008) using: (a) GOSAT low-earth orbit observations (GOSAT_LEO); (b) GEO-CAPE observations with a TES-like instrument (TES_GEO); (c) GEO-CAPE observations with a GOSAT-like instrument (GOSAT_GEO); (d) GEO-CAPE observations with a multi-spectral instrument (MULTI_GEO). Zero values correspond to emissions with no constraints from observations, while values of one correspond to emissions entirely constrained by observations. The DOF for each inversion, which is the sum of all diagonal elements of the model resolution matrix, is also indicated.

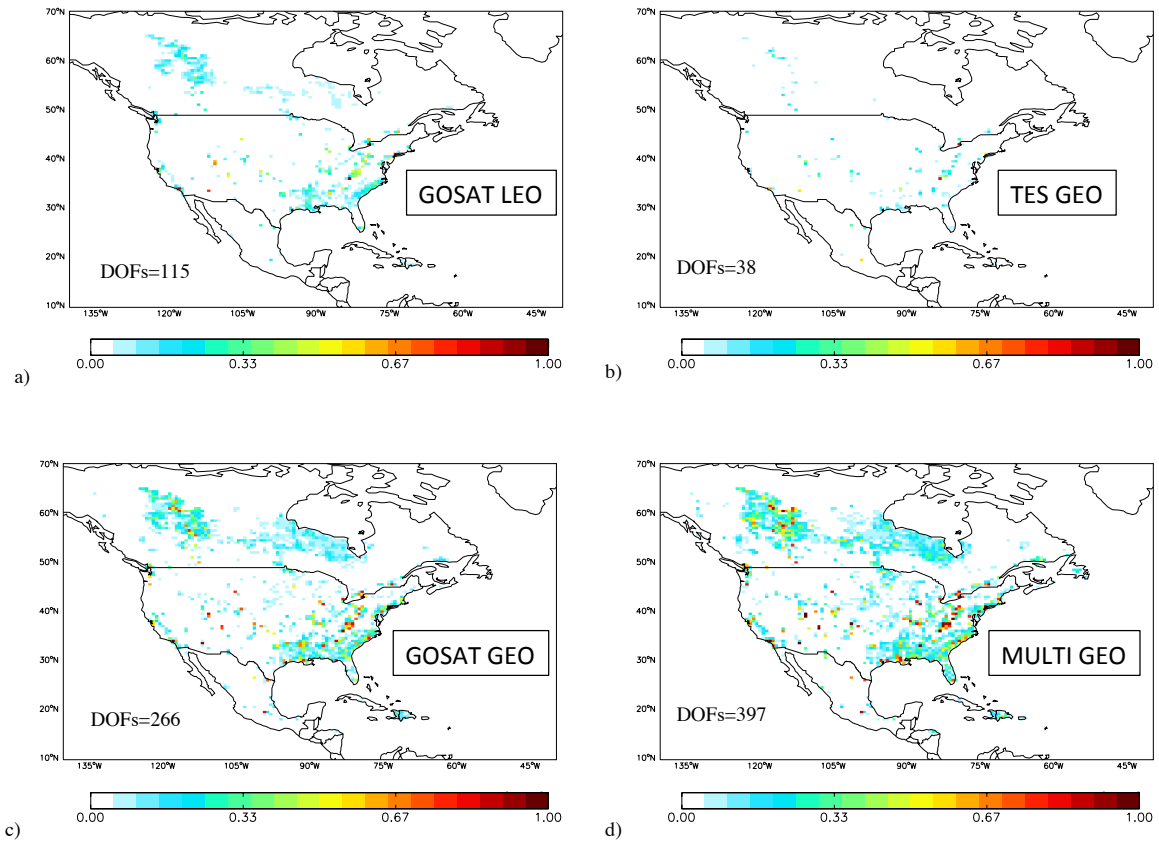


Figure 6. Relative error variance reduction for a 3 day methane emission optimization (1–3 July 2008) using: **(a)** GOSAT low-earth orbit observations (GOSAT_LEO); **(b)** GEO-CAPE observations with a TES-like instrument (TES_GEO); **(c)** GEO-CAPE observations with a GOSAT-like instrument (GOSAT_GEO); **(d)** GEO-CAPE observations with a multi-spectral instrument (MULTI_GEO). Zero values correspond to emissions with no constraints from observations, while values of one correspond to emissions entirely constrained by observations. The DOF for each inversion, which is the sum of all diagonal elements of the model resolution matrix, is also indicated.

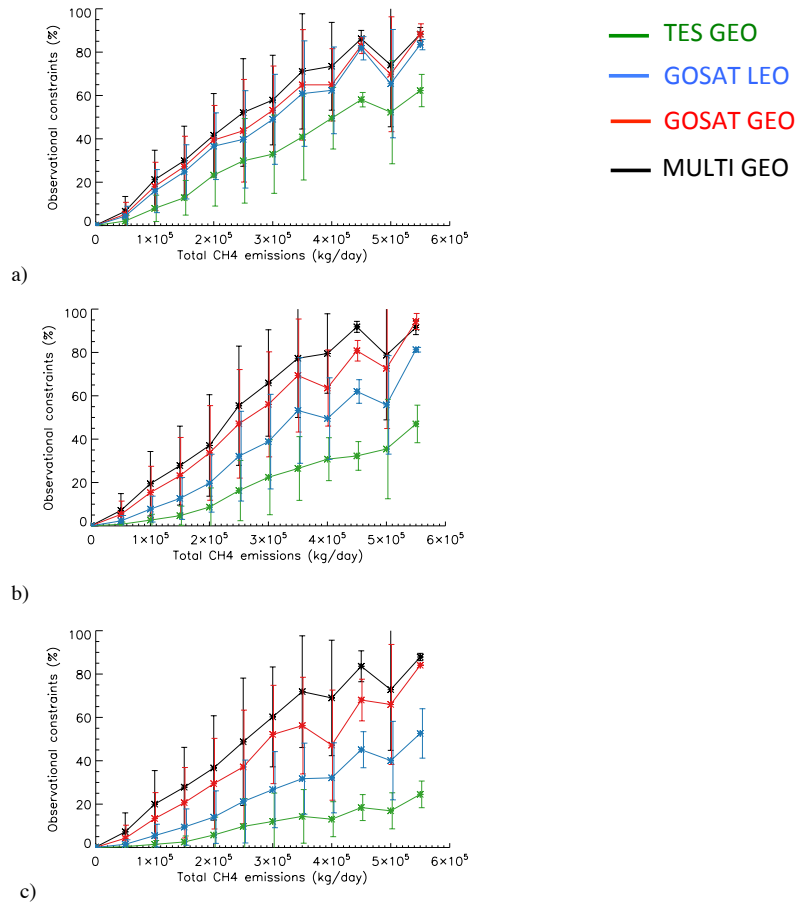


Figure 7. Relative error variance reduction as a function of methane emission magnitude for: **(a)** a 30 day (1–30 July 2008); **(b)** a 7 day (1–8 July 2008); and **(c)** a 3 day (1–4 July 2008) inversion. Blue: GOSAT low-earth orbit observations (GOSAT_LEO); green: GEO-CAPE observations with a TES-like instrument (TES_GEO); red: GEO-CAPE observations with a GOSAT-like instrument (GOSAT_GEO); black: GEO-CAPE observations with a multi-spectral instrument (MULTI_GEO). Results for a 3 day MULTI_GEO inversion are also shown in purple (top). The vertical bars indicate the standard deviation of observational constraints within each bin.

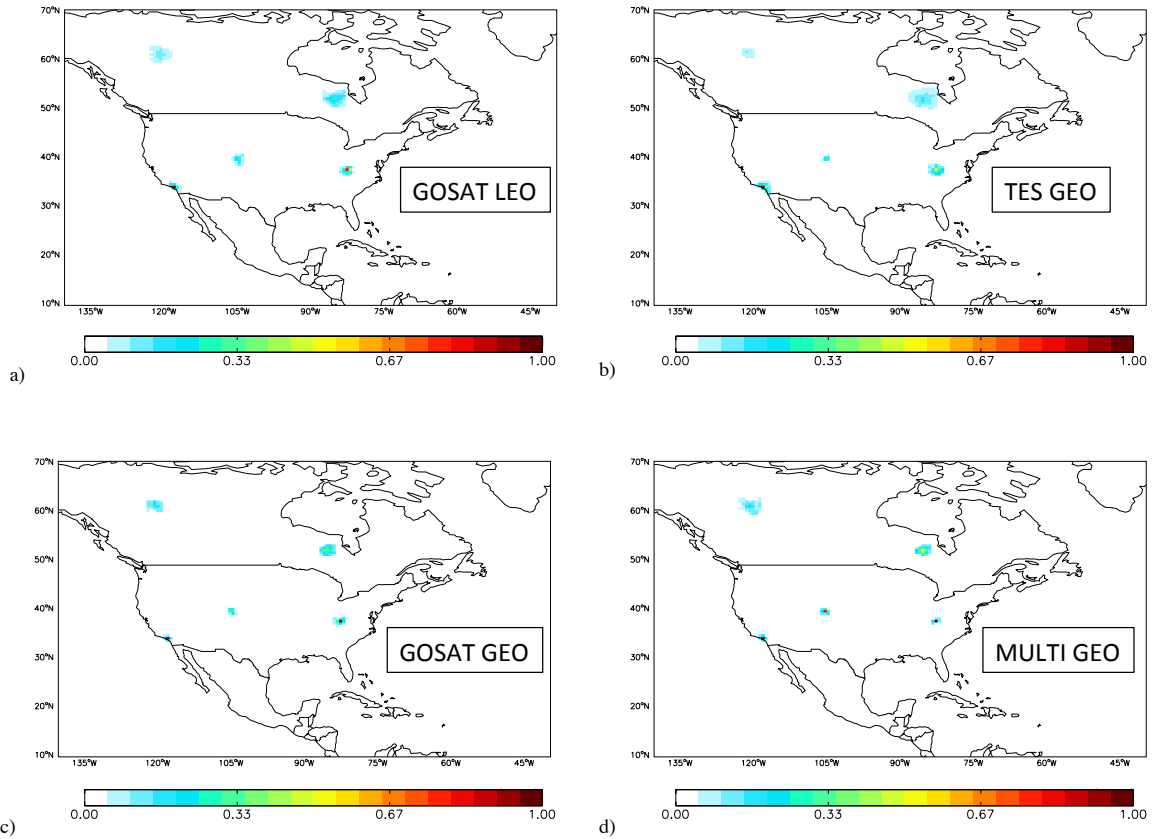


Figure 8. Rows of the model resolution matrix (unitless) for five locations for a 7 day inversion (1–8 July 2008), using: **(a)** GOSAT low-earth orbit observations (GOSAT_LEO); **(b)** GEO-CAPE observations with a TES-like instrument (TES_GEO); **(c)** GEO-CAPE observations with a GOSAT-like instrument (GOSAT_GEO); **(d)** GEO-CAPE observations with a multi-spectral instrument (MULTI_GEO). Coordinates of the five locations considered are reported in Table 1 and approximately correspond to the peak value of each structure on the maps.

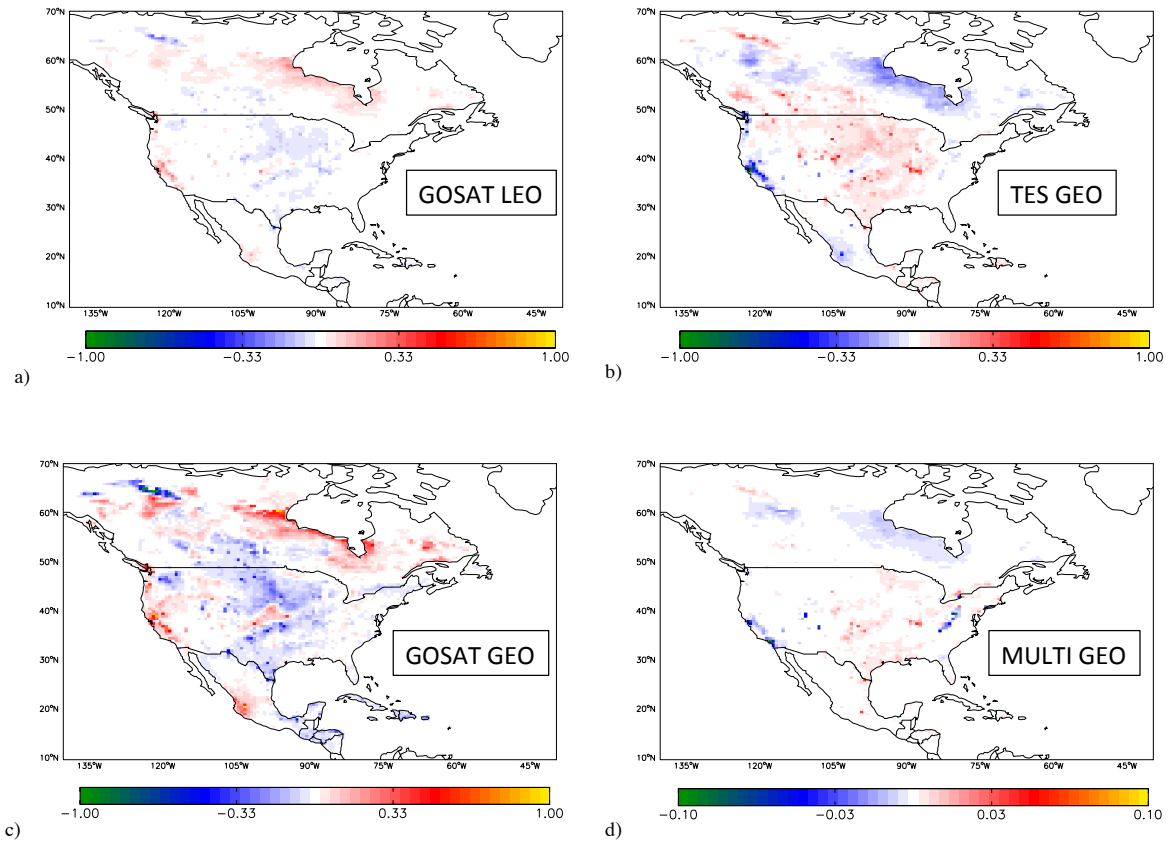


Figure 9. Sensitivity of the optimized emission scaling factors to uncertainties in boundary conditions for a 7 day inversion (1–8 July 2008), using: (a) GOSAT low-earth orbit observations (GOSAT_LEO); (b) GEO-CAPE observations with a TES-like instrument (TES_GEO); (c) GEO-CAPE observations with a GOSAT-like instrument (GOSAT_GEO); (d) GEO-CAPE observations with a multi-spectral instrument (MULTI_GEO). Shown is the impact of perturbations of the boundary condition concentrations with gaussian distribution $\mathcal{N}(0,16 \text{ ppb})$ on the optimized scaling factors. Note the different color scale for the MULTI_GEO configuration.

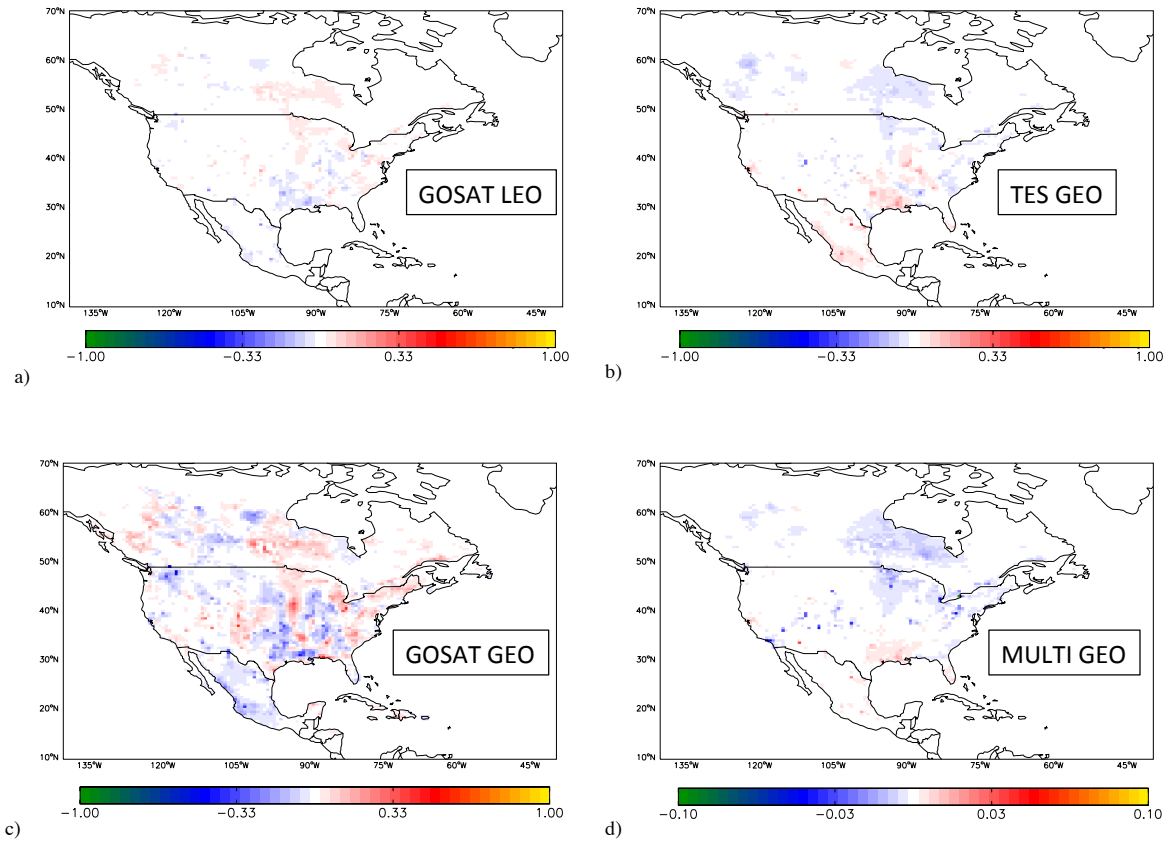


Figure 10. Sensitivity of the optimized emission scaling factors to uncertainties in initial condition concentrations for a 7 day inversion (1–8 July 2008), using: **(a)** GOSAT low-earth orbit observations (GOSAT_LEO); **(b)** GEO-CAPE observations with a TES-like instrument (TES_GEO); **(c)** GEO-CAPE observations with a GOSAT-like instrument (GOSAT_GEO); **(d)** GEO-CAPE observations with a multi-spectral instrument (MULTI_GEO). Shown is the impact on the optimized emission scaling factors of perturbations of the boundary layer and free troposphere initial CH_4 concentrations with gaussian distributions $\mathcal{N}(0,22 \text{ ppb})$ and $\mathcal{N}(0,46 \text{ ppb})$, respectively. Note the different color scale for the MULTI_GEO configuration.