Below are concerns of each reviewer in italics, followed by the author's response and any actions that were taken to resolve it. At the end is a markup of the revised paper.

Response to Reviewer 1:

Minor comments

line 109-112: what are the range of estimates for these numbers? You provided them for the Karion studies, but not these other two.

Added the ranges of each to the paper.

line 184: Perhaps cite the latest EPA greenhouse gas inventory

We cited one from the previous year here because that was where the emission factor came from when this project was active. We cannot cite the most recent report here as the emission factors change from year to year and so the number may be different from the one we used.

line 260-261: do these CH4 sinks account for other reactive species in the atmosphere? If not, how would background or elevated levels of ethane or other gases affect these CH4 losses?

This is a fair point. Gases other than CH, are emitted from wells and could possibly decrease the reaction rate of OH with CH, However, the purpose of this section is just to show that even in above average OH concentrations, the CH, sink is negligible. Unless the other gases released from well emissions lead to the production of additional OH, any gas released that will react with OH will only lower the sink's impact further.

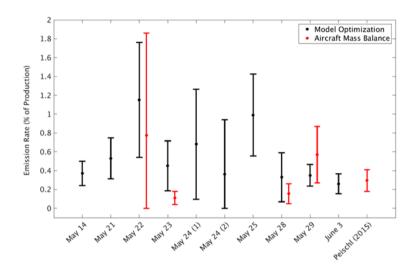
Figure 2: if possible, it would be helpful to know where Bradford County was on this map

Updated the figure with the county highlighted in yellow



Figure 19: I would add the mean model optimization and aircraft mass balance emission rates to this graph, as well as the Omara and Peischl studies, along with their uncertainties. It seems like one advantage this study has, as mentioned in the introduction, is a narrowing of the uncertainties of these previous studies.

I did not add the mean for each as it made the plot a bit heavy and I want to emphasize the spread for each day. I also did not add Omara's number as it comes from emissions only from well production (no gathering emissions) and all wells from that study were measured in the southwest Marcellus, which has a much different composition compared to the northeast Marcellus. However, I did add the number from the Peischl study to the plot, as his study is a single mass balance flight in the region and so it fits in perfectly with the rest of the graph's content. I am not sure our final findings narrowed the uncertainties as intended so much as it discovered how uncertain the previous estimate may have been.



Technical Comments

The author numbering is not in order.

Fixed

line 48: Methane also has an atmospheric sink from chlorine chemistry.

Clarified the sentence to imply the sink was mostly from OH.

line 58: is "mining" the proper word for natural gas? It may be, I just can't say I've heard that before.

Forget you heard it. It is not a good use of the word. Switched to "drilling and transportation of natural gas".

line 64: add "of" to say "quality and quantity of its emission factors".

Added.

line 107: put "per day" before the abbreviation "MMSCFD", instead of after it.

Switched the order.

Response to Reviewer 2

Minor comments are addressed first. Major comments which have not been addressed through fixes in the minor comments are then addressed afterwards.

Many Minor Comments:

L51: Shouldn't you rather say that for emission rates larger than 3 %? Otherwise the sentence does not define a threshold.

Changed to "greater than"

L232-238: This paragraph should be directly in front of the one describing the data assimilation strategy finally used in the study (L246-253).

Moved the paragraph accordingly

L261: "The important time scale for the mass balance approach is not alone the flight duration but also the transport time from the upwind to the downwind interface of the box. Also the time from emission to sample should be discussed, as this is the crucial one for the inverse modelling."

This segment is discussing how oxidation can decrease the background concentration of CH, during the duration of the flight. The emissions should not matter in this calculation because OH would be reacting with the entire CH, mole fraction (i.e., the background + enhancement), not just any enhancement in the region (a 40 ppb enhancement would change the CH4 mole fraction by 2%, so changes to the reaction rate would be minimal). So it will be affecting all parts of the flight almost equally throughout the time of the flight and shouldn't have any relation to the distance the emissions are travelling from the well to the downwind transect.

However, I do agree that it is good to know the time it takes for emissions to travel to the downwind transect, if only to better understand how representative the upwind transect is of the air mass measured by the downwind transect. Given the range in wind speeds, it takes about 2-4 hours. This is mentioned briefly in the error analysis section.

L312 and elsewhere: I don't like the terminology here. "observed CH4 enhancement associated with upstream natural gas". The described quantity contains information from both model and observation and, thus, is not purely 'observed'. Maybe the term 'observation-based' or something else that identifies the real character of the data could be used instead.

The term "observation-derived" replaces now "observed" across the entire document.

L321: How frequent and how large are negative values of X_GasO?

Added sentence in paper describing negative values. 16% of the observation-derived enhancements are negative, but less than 3% are negative by 5 or more ppb.

L352: "Were individual atmospheric densities for each flight neglected? Why?"

They were not calculated, because a +-10 change in the temperature produces a difference in the value (and thus the end result) of 3%. It's a similar change with a +-30 hPa change in the pressure. For 10 flights during a 3 week period with no huge meteorological anomalies, calculating the exact density of the air in the ABL (which will have its own errors) based on a few vertical transects seemed unnecessary.

L416: If this is a good correlation coefficient, what is the range obtained for the other flights? Could this information be added to Table 4 or 5. It could also be used as part of the uncertainty assessment..

Correlation Coefficients for each flight have been added in a column under Table 5

L418f: "Be a bit more specific concerning the near zero differences between model and observation. For the PBL height this statement may be true, however, it is also clear that the model is not able to capture the very sharp increase in potential temperature. I also wonder why the two profiles are shown separately, would be nice to have them on top of each other. Concerning the wind there is a directional shift of about +10 degree in the model. You may call this small, but for the kind of plumes simulated here it might be of importance and should be mentioned."

I do not discuss the gradient of the potential temperature inversion or the offset in temperature between the model and observations because these factors do not impact the physics of the plume. For now, I would like to keep the two plots separated, as the important thing to emphasize with these plots are the ABL height similarities between the model and the observations which can be seen just fine with the plots as they currently are. Combining the plots would draw the reader's attention away from the ABL height and more towards the 1 K difference in temperature, which is not important to this discussion.

Changed L419-20 to reflect a difference in wind direction of 10 degrees.

L424ff: Please add a number for the average and maximum enhancement due to upstream natural gas sources at the northern transect to this discussion. From Figure 9 I would think these numbers are around 30 and 80 ppb. It puts the other contributions into perspective.

Adjusted line in the paper to give a range of the downwind enhancement.

L473ff: This discussion on the uncertainties due to shifts in the simulated wind direction is very hard to follow. Figure 16 does not speak for itself. You will need to explain in more detail what was done to derive the figure. This discussion should actually given when the cost function is introduced.

Moved discussion of optimization technique and how it addresses transport uncertainties to methods section. Removed Figure 16 and wrote better explanation as to why sum of squares was not used as optimization technique. Simplified paragraph and associated figure is provided below.

"The decision to use a scalar cost function rather than the sum of squares is to account for possible misalignment between any observed CH4 plume and modelled plumes. There are two potential ways in which misalignment may occur. One possibility is that the modelled wind direction differs from the true wind direction, leading to a plume in the model that is off-centre in relation to the observed plume. The other possibility relates to how the model treats emissions from natural gas as a uniform percent of production. In reality the emissions are more random in nature, and thus the plume may not always develop over the wells with the largest production values. If a cost function is used that minimizes the sum of the squares, any misalignment between the modelled and observed plume will result in the peak of the modelled plume aligning with the height of the tail of the observed plume (Figure 5). Unless the observed plume aligns perfectly with the modelled plume, the optimized emission rate using a sum of squares approach will always bias low. By using a scalar cost function, we solve for an optimized emission rate that results in a plume with the same area under the curve compared to the observed plume (Figure 5). This methodology is not impacted by any misalignment between the modelled vs. observed plumes, preventing the low biases associated with a sum of squares minimization."

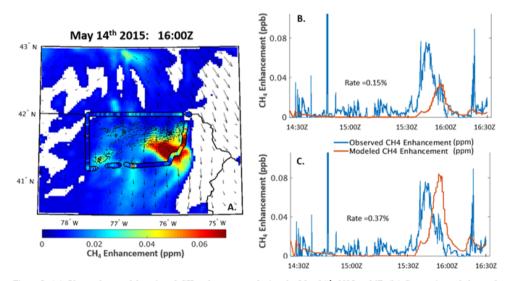


Figure 5: (a.) Observed vs model projected CH₄ enhancements during the May 14th, 2015 at 16Z. (b.) Comparison of observed natural gas enhancement to modelled natural gas enhancement along flight path, with upstream emission rate optimized by minimizing the absolute error between the datasets. (c.) Same as previous, but optimized by minimizing the sum of the error between the datasets.

L486: Should it be wind speed instead of wind direction?

It should be wind speed. Changed.

L498ff: "The way the wind speed error is assessed biases could be introduced. The average over all wind observations by the aircraft does not necessarily reflect the average wind speed within the boundary layer, since the sampling is not uniform (especially with height). It seems calculating an average vertical profile of wind speed from all the observations and comparing that to an average model profile generated from the same sampling locations should be more robust. Was this considered and decided that the aircraft sampling was sufficiently uniform?"

Most of the time the aircraft is flying at a constant altitude. We have about 2-3 vertical profiles (mostly spirals) from the aircraft for most flights, but wind measurements can have large errors while the aircraft rises/falls and turns, both of which occur during a vertical spiral. So we cannot accurately construct a vertical profile of the winds in the ABL using the aircraft data. We could compare the model to surface station measurements in the area, but surface data poorly reflects on model inaccuracies within the boundary layer. Comparing the model directly to the aircraft observations when it is flying steady in the boundary layer is the best method we have of understanding the model's performance throughout the area, and also will capture all of the potential spatial variability as well. I do acknowledge though that due to a lack of wind measurements with height, if the model has a bias that is not uniform with height this will not be captured in the error analysis.

L506: "There are more advanced techniques for estimating boundary layer heights from model and observational data than just 'looking' at the potential temperature. For example Bulk Richardson methods are quite useful in situations when potential temperature alone is not providing explicit results. The example shown here is an easy case but I wonder about the other situations mentioned in the text and especially how large the uncertainty of the estimates becomes in such situations."

Reading the sentence associated with this comment, I may have implied more uncertainty than there actually was. For all 10 flights, the potential temperature inversion could clearly be spotted in the model. This is also true for 9 of the aircraft flights. Only on one day (May 22nd, 2015) did the observational data not clearly show a large potential temperature gradient until ~3000m, with a smaller wiggle in the profile a bit lower. For this one though you could see a quick shift in the all of the trace gases at that wiggle, and so we could conclude that this represented the mixing height. We did not go into a more complicated methodology for deriving the ABL height because we felt it was unnecessary for our cases. Had the potential profiles been messier and vague, we would have then resorted to more complex methods of calculating the ABL height.

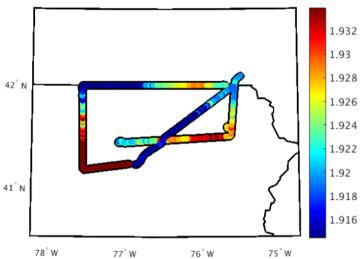
I have revised the sentence in the manuscript to make it more clear that there was little ambiguity in finding the ABL height.

L542f: "How was this done exactly? Did you simply run 3 cases with 1) reference background, 2) background -5 ppb, 3) background +5 ppb? Or did you do a Monte Carlo approach where the +/- 5 ppb could be understood as the standard deviation of a normal distribution? Please clarify and justify the approach also in the light of possible non-linear effects that would not be covered by simply running 3 cases."

It was done by just running 3 cases to establish the limits because the effects changing the background has on the solution is nearly linear. Unlike using a sum of squares approach in our cost function, the cost function we use tries to minimize the difference between the area under the curve for the observations vs the model. Another way to think of it is that we are minimizing the total enhancement between the sum of the observation-derived enhancements and the sum of the modelled enhancements. By decreasing the background by 1 ppb, we are adding (1 ppb * # of observations) to the total enhancement. By decreasing the background by 2 ppb, we are adding 2 ppb * # of observations) to the total enhancement. The increase (or decrease) is linear, as is the effect on the cost function. The only effect that is non-linear occurs in the process the zeroing of any negative observation-based enhancements (of which you will have more of if you increase the background by 5 ppb), but the impact of this is small on the symmetry of the error bar. So a Monte Carlo approach was not needed due to the simple nature of the errors in this section.

L548f: It is mentioned that the +/- 5 ppb assumption for the background uncertainty can have different impacts depending on the magnitude of the observed plume and this is related to wind speeds and PBL heights. How can you justify the use of a constant value for the background uncertainty? Shouldn't this also change with meteorological conditions? For example it is mentioned earlier that it will be large when there is more entrainment, which is usually the case while the PBL is still developing, and when plumes from other sources are advected. Wouldn't it make sense to take this variability in the background uncertainty into account?

In truth, a lot of the changes we see in the background we just don't understand. Below are the methane observations from within the ABL for our flight on May 22nd.



This was a 3 hour flight between 1:30 pm-4:30pm local time. Winds were steady on this day and the boundary layer was deep throughout the duration of the flight. We should expect minimal changes in

the background in these conditions. But near the very beginning of the flight (bottom left), values were their largest despite being away from most sources. Meanwhile, values near the end of the flight suddenly drop off by 10 ppb despite having flown over the same area twice just one hour earlier. This is the most extreme example I can present of the randomness of the background variability, but in my experience with aircraft data, there are many instances in which sudden changes in the background can be observed which appear to be unrelated to entrainment or a simple calculation in how much the boundary layer changed. Much of the differences we see in the background values may have to do with heterogeneity of the large CH₄ plumes which develop regionally across the U.S.. As a curiosity, we ran WRF-Chem for 1 month at 27 km resolution using the EPA U.S. CH₄ emissions inventory (Maakkassers *et al.*, 2016) as input and found that the average difference in enhancement between two locations 71 km away in our study region was 5 ppb. These modelled differences are caused by the spatial gradients that naturally develop in the methane concentration field due to the transport of methane emissions across the central and eastern U.S.

In short, we picked a background variability of ±5 ppb to represent a conservative estimate of how much variation we could see in a flight. We did not believe that any calculation we could perform with the data we had could sufficiently describe our uncertainty with the background value estimate, so we chose a range that seemed large without being unreasonable.

L557f: "Again the question: is [the non-natural gas emissions adjustment] done by running by simply running 3 cases or in a Monte Carlo fashion. Did you scale all categories with the same factor at the same time or test combinations as well? How do you obtain symmetric uncertainties in the final emission rate when you use asymmetric uncertainties in the non-upstream natural gas emissions?"

The 50-200% range was a typo and should have read 0-200%. However, we agree that the initial error analysis from the inventory could be improved upon. To improve this section, we followed the advice provided and ran a Monte Carlo simulation for 10,000 iterations, selecting a multiplier from a uniform distribution ranging from 0 to 2 for each of the 8 non-upstream gas sources. This allowed for a much more realistic understanding of the errors compared to assuming all sources were incorrect by the same multiplier. The resulting patterns for each day were close to Gaussian. For each day, we fit a Gaussian and took the 2 sigma range as the uncertainty. Most days did not see much of a change from the previous uncertainty assessment, but days where the coal plume interacted with the observations (May 24th May 25th) saw an increase in the error. The methodology for this has been updated in the paper.

L505, L562: "Table 5 should already be referenced here and not only at the very end of the section."

Added for L562. I didn't add a reference at L505. That's describing a different methodology related to Table 4. I did add another reference to both Table 4.5 when the error analysis is first introduced.

L571: "The estimator for the model performance uncertainty is a normalised root mean square error (NRMS) and it should be called like this to reduce the confusion. However, I am not convinced that this is the best estimator for what you want to achieve. One problem is the use of the optimised simulation, which means you have already used the information contained in the observations and thus the uncertainty estimator is not independent anymore. Furthermore, NRMS is not a good measure of what you call "similarity of pattern". The Pearson correlation coefficient should be better suited for this

purpose. Please justify your choice. If you stay with a RMS estimation it should be applied to the prior simulations and the bias should be removed before the RMS is estimated."

We had originally looked at the correlation as a way to quantify uncertainty, but we found there were some interesting cases where the correlation did not accurately describe problems with the model vs. obs comparison. For a great example, look at Figure 17 (May 22nd Loop1 vs May 28 Loop2).

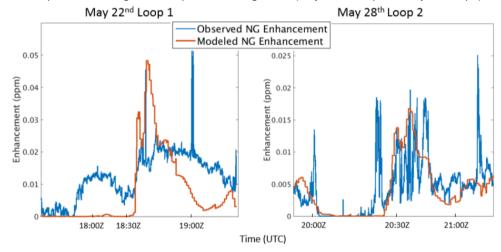


Figure 17: Comparison of observed natural gas enhancement to modelled natural gas enhancement for segments along the (left) May 22nd flight and (right) May 28th flight. A distinct lack of representativeness of the observations in the modelled enhancement can be seen in the May 22nd flight compared to the May 28th flight.

Anyone looking at the figures would immediately conclude that the observations on May 28th were modelled better than May 22nd. However, the May 22nd Loop1 flight actually has a correlation coefficient of 0.58 while the May 28th Loop2 flight has a correlation coefficient of 0.57. Of all the methods we tried, we found the NRMS method we use resulted in the error estimates that matched with a common-sense approach.

I do not fully understand the request to apply our methodology to the prior. Unlike an atmospheric inversion, our forward-based modeling experiment does not have a true prior. The number we selected for our first-guess emission rate (0.13% of production) has 0 bearing on the final results. It only provides the model with the spatial pattern the plume will take, and is scaled entirely by the observations. Had we started with a first-guess emission rate of 82,350% of production, the final result and uncertainties would not change.

Added a line addressing that the formula used is a modified NRMS to avoid confusion.

"L579f: You also assume Gaussian uncertainty distributions, which seems oversimpli-fied."

As addressed above, the certainty of our background uncertainty parameter is very uncertain. The Monte Carlo for the non-natural gas emissions uncertainty produced mostly Gaussian-shaped distributions, but this was assuming an arbitrary range of variations in the emissions of each source which may also have spatial variabilities that we don't understand, and the model performance error is just a method we picked to try and best match our own understanding of what the errors associated with mismatch should look like. Assuming Gaussian uncertainty distributions is probably a stretch, but there was no methodology we could come up with where we would feel any more confident in our final error bounds.

To counteract the uncertainty in our uncertainty, it should be noted that we tried to be very conservative with our uncertainty parameters. A ± 5 ppb variation with the background is large. A 0-200% range in the emissions from non-natural gas sources is large. We did not want to underestimate the uncertainty, as is often done, so we tried to be as fair with our analysis as possible to produce a 2σ range which would contain the true emission rate 95 times out of 100.

L645:" Figure 18 can go into a supplement without any loss of information. Instead give the rate of change of the background with time as a number in the text."

Added the rate of change in the text, and Figure 18 is in consideration to be moved to supplemental (see major comments below)

L652: "Does this mean the final uncertainty estimate is the mean over the uncertainties for each flight?"

From that sentence, we use the standard error from the four flights to come up with the uncertainty range.

L663ff and Fig. 19: "The variability in emission rate estimates for individual days as shown in the figure are not discussed in any detail.

This is a good point. Added a segment in the early part of the discussion section to discuss reasons for variability in the emission rate.

Of special interest is the question of agreement between the two methods as it could lend some additional trust in the applied methods. However, the figure seems to raise more questions in the reliability. For example if 29 May is the golden day (as described earlier in the text), why are estimates from both methods so different?

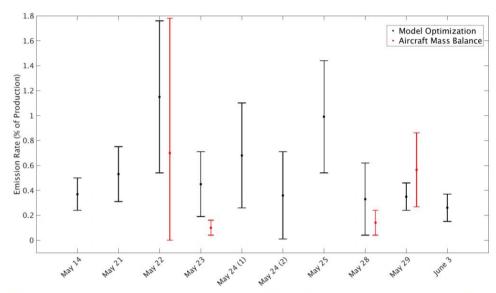


Figure 19: Calculated upstream natural gas emission rates using (black) model optimization technique and (red) aircraft mass balance technique.

The values on May 29th from the aircraft mass balance and model optimization are 0.57% and 0.35%, differing by 0.22% of production. They even fall within each other's error bars. Claiming the two values are "so different", is odd, and should not be used to question the reliability of the model optimization method. Furthermore, the aircraft mass balance method has sources of error which differ from errors associated with the model optimization technique. A perfect match between two methods each with their own unique sources of error is unlikely.

I do discuss briefly in the paper that the relative closeness of the results in these two methods gives credence to the model optimization technique. But that is not to say the aircraft mass balance method is the golden standard in the field. If the mass balance approach were flawless, bottom-up and top-down emission estimates would be in agreement at this point and there would be little need for this paper.

Also there seems to be no agreement in the variability for the 4 common days. How could you explain the large variability from the point of the emission processes? Is there any dependency on the total production on these days (following the line of thought used later for the discussion of different basins)? Please speculate on this otherwise the reader is left with the impression that your given uncertainty estimate is rather optimistic."

The topic of emissions variability is briefly touched upon in the introduction, but I've added a segment in the discussions to re-address some of the possible reasons for the variability in the totals. These reasons include the nature of natural gas releases from stochastic events such venting, flaring, liquids

unloading, ect., as well as the possibility that the days with large emission rates also have large uncertainties due to the complex scenarios from those days.

L666f: Repeat the range of emission rates from other studies here.

Added a line addressing this rate compared to other regions.

L696ff: The flight described in this section and shown in Fig. 21 is not part of the main analysis of the paper and should be omitted as it does not provide any additional insights and aspects that have not or could not be shown on the other flight examples.

On the contrary, this flight is the perfect example to demonstrate why it is so advantageous to model the plumes with a direct model instead of the backward adjoint model. Until this study, a mass balance from a single flight was the only top-down study performed in the northeastern Marcellus. There was little knowledge of how the coal plumes 500 km away would interact with the flight in this region. By running a transport model, we're able to see that the background of this flight is compromised by the heterogeneous nature of the coal plume, thus explaining the complex signals observed by the aircraft (it should be noted that the highest CH₄ values observed in *Peischl et. al.* 2015 were **upwind of the wells)**. Even if someone does not plan to use the transport model to solve for the emissions in a region, it can still be useful to better understand the complexity of background plumes on a given day and decide whether a mass balance would be appropriate to use. By using the flight from *Peischl et. al.* (2015), we not only show the usefulness of using a transport model for future studies, but also emphasize the potential need to relook at flights from earlier studies and check for whether complex plume structures may be adding additional uncertainty to the regional emissions estimate.

Addressing the major concerns directly

Model optimisation technique (inverse modelling): The applied optimisation technique uses a very simple cost function to find an optimal factor between 'prior' and 'posterior' emission rates. Why did you not apply a Bayesian approach that could take uncertainties in the prior values and observations/simulations into account? Much of the following uncertainty assessment could then be used in the optimisation step and also more reliably give posterior uncertainties. Also, why do you not use sum of squares in the cost function as is commonly done? Some explanation is given later in the text, but without further discussion of Figure 16 this remains useless. Please discuss why your method should provide more reliable results as other similar studies that have used a Bayesian framework.

As far as I'm aware, there are not any similar regional methane emission studies that have combined a Bayesian framework with an aircraft campaign. We know of few ongoing studies but to date, none of them have been published. Few top down studies used gas-to-gas ratios and backward footprints to estimate point source CO₂ emissions (e.g. Brioude et al., 2012) but without any prior emissions. One of the advantages of a Bayesian methodology is that it allows one to use and propagate their errors from a prior guess to the posterior estimate. However, it is for that same reason that a Bayesian approach would be inappropriate here. We have little information regarding the errors associated with this study. The background CH₄ values, the emissions inventory, and the model transport error are all poorly understood. This includes the 'prior' emissions estimate of natural gas as well, which is just the median

emission rate from 17 UNG wells in southwestern PA. Any Bayesian approach without reliable uncertainty estimates will not produce reliable results (Chevallier et al., 2006).

One of the greatest advantage to our forward-based modelling approach is that you are able to visualize plumes and partition the problem into components, directly observing the uncertainties associated with the optimization process. The forward modelling approach lets you directly see the structures of the different plumes on a given day and how they compare to observations. This makes it simple to know your problems for a specific day. If there are far-reaching plumes influencing the observations, you know to be cautious with the emissions estimate. If plumes appear to consistently misalign slightly between the observations and the model, you know there's likely a transport issue. If observations shift in regions where no plumes are modelled, you know you are dealing with either a missing source or background variability. And because emissions scale linearly with the resulting plumes, once the model is run it becomes easy to make adjustments to see how much impact changes can have on your results. These different aspects are purely and simply invisible in a backward mode, for which the adjoint only points to the simulated area of influence. The difficulty with the direct approach is to rigorously quantify the final uncertainties, but uncertainties obtained from a Bayesian will be no more reliable if the prior uncertainties are not known.

So while the approach we use in this study may be simplistic, we do not consider that to be a weakness. Rather, the ease-of-use associated with this method allows it to be readily applied to any other study, past, present, or future, so long as an emissions inventory can be compiled for the region.

We have responded to the "sum of squares" issue under minor comments. We have considered methods such as Nearest Neighbor Search (NNS) but found it tedious and unsatisfactory as it would look for surrounding pixels and would therefore require an adjoint model.

Uncertainty assessment of obtained emission rates: The uncertainty assessment of the obtained emission rates for both the 'model optimisation' as well as the mass balance method contains a lot of arbitrary assumptions and does not seem to be statistically sound. Their seems to be an overall assumption of Gausssian uncertainties (although not explicitly mentioned), which does not seem to be justified for several of the sources of uncertainty. Also the way the uncertainties are propagated through the individual methods remains very vague and could be improved by using a Monte Carlo type uncertainty assessment. These concerns are repeated in detail below.

We addressed this concern at multiple points under minor comments section. To summarize here, we have propagated errors when possible in the revised manuscript (non-Natural Gas sources), and found that our Gaussian assumption matches closely the exact error distribution in this case. For the other error terms, we have statistical information on transport errors (from other studies at similar scales, e.g. Deng et al., 2017) to confirm it, or reasons to use a Gaussian propagation of the standard deviations when the distribution remains unknown (e.g. prior errors on emissions). We also want to emphasize that the absence of known prior uncertainties for emissions will significantly impair our ability to diagnose the full error distributions with a Monte Carlo approach, Therefore, random sampling such as Monte Carlo would not solve the problem of unknown error distributions. In general, we have been very conservative in our estimates, using the high end of the error values. Unless more information on error sources becomes available, we think that propagating errors under Gaussian assumptions remains valid and perfectly justified.

Length: With 21 figures, 8 tables and a total of 50 draft pages the manuscript is quite lengthy and it would profit from shortening and restructuring. Below I suggest a number of figures that could easily be omitted or moved into a supplement without loss of information. Furthermore, I strongly encourage to incorporate the description of the uncertainty assessment method, which is now given within the results section, into the methods section along with or following the description of emission rate estimation itself. In this was many repetitions could be avoided and the paper could really focus onto the results in section 3.

A major restructure was done to move uncertainty assessment into methods section and eliminate duplications associated with having those split previously. Results are now concise and focused only on results, rather than methods. The number of figures has been reduced by combining Figures 1 and 20, and the removal of Figure 16. If the editor feels that the length of the paper is an issue, further changes can be made, such as moving Figure 4 and 18 to supplemental as well as the entire description of the mass balance method used in this paper. However, if the editor feels that length is not an issue, we would prefer to keep these sections in the paper to retain its flow.

Technical Fixes

L26: Abbreviation (WRF-Chem) not introduced before. Do so on line 25. Added abbreviation

L48: "transforms" instead of "transform"

Due to interactions... which transform. Grammar is correct

L59: Add "United States" in front of "Environmental Protection Agency". Added

L348: Why are the terms U, D, and L given in braces? Not necessary. Braces removed.

L496: "as long as" instead of "so long as". Changed

Fig. 7: Do not show wind arrows in center panel. Not visible anyway and one gets the general flow direction from the shape of the plumes. Show center panel with same size as the individual tracers. Also use exactly the same color scale for all sub-panels. It looks like the center plot does have fewer color levels than the others. Each panel (also the center panel) requires a label.

We respectfully disagree with this comment and prefer to keep the wind arrows to illustrate the direction of the flow for this very specific case. The coal plume coming from far away is explained by the regional circulation. The arrows remain small and should not decrease the readability of the figure. We modified the colorscale to match better with individual tracer figures. Did not resize middle figure to match other figures. Middle figure is most important and thus should be drawn as such. I think the label being referred to each panel needing was lat/lon. Added to the main figure. Not added to smaller

figures as it makes things messy and unreadable. I would think the reader would have a good understanding of where they are in space using middle figure as reference.

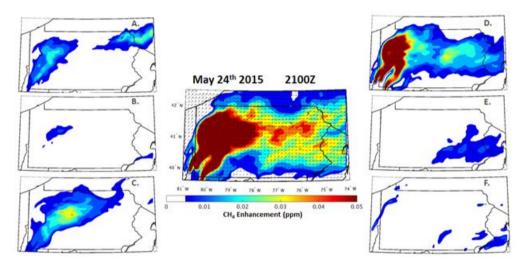


Fig. 11: Show observed and simulated profile on top of each other (see also comment above).

The purpose of the figure is to show an example of a theta profile for both the observations and the model and how the abl was found for each. The current figure accomplishes this fine.

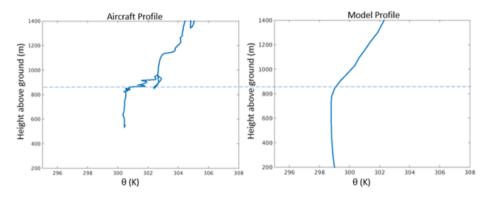


Fig. 12 and others: There is no mentioning of the wind arrows in the figure legend. Are these near surface or mean PBL winds? There is also no reference vector that would allow inferring

the wind speeds from the arrow length. Choose a different color for the wind vectors. They are not well visible in the righthand figure.

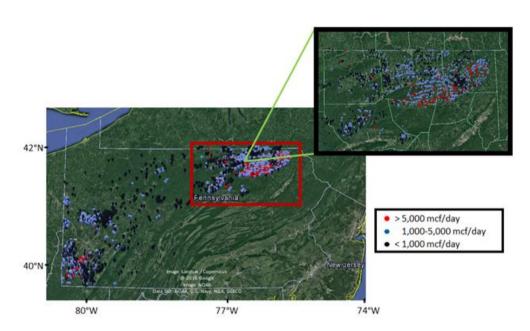
Added description of height level of model used in figure. Wind speed by arrow length changes in a complex manner from figure to figure with no easy way to retrieve its scale. Given the context of the figure, the reader will likely not be interested in small changes in the wind speed but rather the wind direction. If a reader wants more information on the wind speed, he can reference Table 3 which contains information on the mean wind speed for each flight. If the editor feels that a wind speed legend is necessary for these figures, then they will be added in. In terms of wind arrow color, it is difficult to pick a color that works everywhere due to the various colors on the plots associated with the plumes. For figure 12 this should not be an issue as it is a two panel figure where wind directions are clearly visible on the left panel. In figure 14 the wind direction is consistent throughout the domain and should not be too difficult for the reader to extrapolate the information.

Fig. 19: Mention in the caption what error bars represent.

Added description.

Fig. 20: This information could also be merged into Fig. 1 and save the additional figure here.

Merged well production information from Figure 20 into Figure 1. Removed Figure 20 from the paper.



Other Changes

Fixed error in Corrected NG Emission Rate (Table 4: WS/ABL bias table), where errors were miscalculated.

Fixed error in calculation of mean and uncertainty regarding aircraft mass balance emission rate.

Minor text fixes.

Quantifying methane emissions from natural gas production in northeastern Pennsylvania

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- Abstract. Natural gas infrastructure releases methane (CH₄), a potent greenhouse gas, into the atmosphere. The estimated emission rate associated with the production and transportation of natural gas is uncertain, hindering our understanding of its greenhouse footprint. This study presents a new application of inverse methodology for estimating regional emission rates from natural gas production and gathering facilities in northeastern Pennsylvania. An inventory of CH₄ emissions was compiled for major sources in Pennsylvania. This inventory served as input emission data for the Weather Research and Forecasting model with chemistry enabled, (WRF-Chem), and atmospheric CH₄ mole fraction fields were generated at 3km3 km resolution. Simulated atmospheric CH₄ enhancements from WRF-Chem were compared to observations obtained from a three-week flight campaign in May 2015. Modelled enhancements from sources not associated with upstream natural gas processes were assumed constant and known and therefore removed from the optimization procedure, creating a set of
- processes were assumed constant and known and therefore removed from the optimization procedure, creating a set of observed enhancements from natural gas only. Simulated emission rates from unconventional production were then adjusted
- 36 to minimize the mismatch between aircraft observations and model-simulated mole fractions for ten flights. To evaluate the

method, an aircraft mass balance calculation was performed for four flights where conditions permitted its use. Using the model optimization approach, the weighted mean emission rate from unconventional natural gas production and gathering facilities in northeastern Pennsylvania approach is found to be 0.36% of total gas production, with a 2σ confidence interval between 0.27-0.45% of production. Similarly, the mean emission estimates using the aircraft mass balance approach is calculated to be 0.3440% of regional natural gas production, with a 2σ confidence interval between 0.960%-0.6272% of production. These emission rates as a percent of production are lower than rates found in any other basin using a top-down methodology, and may be indicative of some characteristics of the basin that makes sources from the northeastern Marcellus region unique.

1 Introduction

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The advent of hydraulic fracturing and horizontal drilling technology has opened up the potential to access vast reservoirs of natural gas previously inaccessible, shifting energy trends in the United States away from coal and towards natural gas (EIA, 2016b). From a greenhouse gas (GHG) emissions perspective, natural gas has the potential to be a cleaner energy source than coal. For every unit of energy produced, half as much carbon dioxide (CO2) is emitted through the stationary combustion of natural gas in comparison to coal (EPA, 2016). However, during the process of extracting and distributing natural gas a percentage of the overall production escapes into the atmosphere through both planned releases and unintended leaks in infrastructure. Though these emissions may be small from an economic perspective, their climatological impacts are not negligible (Alvarez et al., 2012; Schwietzke et al., 2014). Methane (CH₄), the main component of natural gas, is a potent greenhouse gas with a global warming potential over a 20-year period (GWP₂₀) of 84 (Myhre et al., 2013). Over a 100-year period the GWP is reduced to 28 due mostly to interactions with the hydroxyl radical which transform the CH₄ molecule to CO2. Depending on which timespan is used, the relative climatological impacts of natural gas as an energy source compared to coal can vary. Using the GWP₂₀ value, it is estimated that a natural gas emission rate of onlygreater than 3% of total gas production would result in a natural gas power plant having a more negative impact on the climate than a coal-powered plant. Using the GWP₁₀₀ value, this emission rate threshold shifts to 10% of production (Schwietzke et al., 2014; Alvarez et al., 2012). Complicating matters further, the future climate impacts associated with an increased availability of natural gas extends well beyond a simple greenhouse gas footprint comparison against coal. Lower fuel prices linked to this new reservoir of energy can change the course of future energy development globally. With many states and countries attempting to find a suitable balance between their energy policies and greenhouse gas footprint, it is important for the scientific community to be able to quantify and monitor natural gas emission rates.

The miningdrilling and transfertransportation of natural gas can be broken down into five stages: production, processing, storage, transmission, and distribution. The <u>United States</u> Environmental Protection Agency (EPA) uses a bottom-up approach to quantify these emissions, estimating emission rates per facility or component (such as a compressor, unit length of pipeline, pneumatic device) or an average emission per event (such as a well completion or liquids unloading).

These "emission factors" are then multiplied by nationwide activity data containing the number of components or events associated with each emission factor, and a total emission rate is produced for the country (EPA, 2015b). This bottom-up approach is a practical methodology for estimating emissions over a large scale but has limitations. A bottom-up inventory depends on the quality and quantity of its emission factors and activity data. Emissions from sources in the natural gas industry can be temporally variable and have a wide range of values depending on a number of factors, such as the quality and age of the device and the gas pressure moving through the component. Furthermore, recent studies have shown that a majority of emissions comes from a small percentage of devices, often referred to as "super-emitters", creating a long-tail distribution of emission sources (Brandt et al., 2014, Omara et al., 2016, Zavala-Araiza et al., 2015, 2017, Frankenberg et al., 2016). These-two factors make it difficult to sample enough devices and adequately describe the mean emission rate, thus allowing for significant representation errors in the emission factors. Because emission factors are required for hundreds of different components, these errors can accumulate and lead to systematic biases in the total emissions estimate.

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One way to compare compliment results based on inadequate sample sizes in the bottom-up approach is to measure the aggregated enhancement in the atmospheric mole fraction at larger scales through a top-down approach. Instead of measuring emissions from individual devices and scaling up, a top-down approach takes atmospheric greenhouse gas concentrations measured downwind of a continent (e.g. Bousquet et al., 2006), a region (e.g. Lauvaux et al., 2008), a city (e.g. White et al., 1976, Mays et al., 2009, Lamb et al., 2016) or a facility (e.g. Ryerson et al., 2001) and uses inverse methodologies to attribute the enhancements to potential sources upwind. One of these methods, the aircraft mass balance technique, has been performed at many different oil and gas fields to characterize natural gas emissions (Petron et al., 2012, Karion et al., 2013, 2015, Peischl et al., 2015, Conley et al., 2016). While this methodology is able to capture surface fluxes over a large region, it remains difficult to attribute the emissions to any individual source (Cambaliza et al., 2014). Any sources from within the flux region that emit CH4 will be measured in the downwind observations and be a part of the aggregated regional enhancement. Atmospheric observations also include other sources of CH₄ unrelated to natural gas, such as anaerobic respiration from landfills and wetlands, enteric fermentation from cattle, anaerobic decomposition of manure, CH₄ seepage from coal mining, and many other smaller sources. If the purpose of the study is to solve for the emissions from the natural gas industry, emissions from all sources unrelated to natural gas must be known and removed from the regional flux estimate. Thus, top-down experiments require an accurate CH4 inventory of the study area and any errors associated with the inventory will propagate into the final emissions estimate. A more advanced technique to separate out non-natural gas sources has been developed using ethane as a tracer for natural gas (Smith et al., 2015). However, such methods may struggle in dry gas basins where smaller ethane to methane ratios within the gas can make the ethane signature more difficult to separate out, or in regions where multiple ethane sources are present. And similar to bottom-up methods, top-down studies fail to address temporal variability, with observations from many of these studies having been collected during a limited number of 2 to 4 hour aircraft flights performed over a period of weeks,

In recent years, both bottom-up and top-down studies have aimed at calculating natural gas emission rates, with bottom-up studies generally finding smaller emission rates than their top-down counterparts (Brandt et al., 2014). The

discrepancy between the results from these two methodologies must be better understood if the true emission rate is to be known. Both the bottom-up and top-down approaches have their own inherent sources of error. For the bottom-up approach, a small sample size could result in the omission of any super-emitters, resulting in a low emissions bias. For the top-down approach, difficulty in attributing the measured enhancements to their correct sources can lead to errors when solving for the emissions of a particular sector.

Top-down emission estimates of individual basins have shown variation in the emission rate across the different basins. An aircraft mass balance performed over the Barnett shale in Texas found an emission rate between 1.3-1.9% of production (Karion et al., 2015), yet a similar mass balance study executed over unconventional wells in Uintah County, Utah, calculated an emission rate between 6.2-11.7% of production (Karion et al., 2013). Differences in regional emission rates can perhaps best be illustrated by recent studies in the Marcellus region. The Marcellus shale gas play is part of the Marcellus geological formation running close to the Appalachian mountain chain from West Virginia to southern New York and contains an estimated 140 billion cubic feet of technically recoverable natural gas (EIA, 2012). Reaching peak production by the end of 2015, the Marcellus is the largest producing shale in the U.S., producing 17,000 million standard cubic feet per day (MMSCFD) per day of natural gas (EIA, 2016a). A bottom-up study measuring emissions from 17 unconventional well-sites in the Marcellus found a median emission rate from the wells of 0.13% of production, but estimated a mean emission rate of between 0.5338-0.86% of production due to the potential presence of super-emitters which would skew the mean emission rate towards values higher than the median (Omara et al., 2016). An aircraft mass balance study over northeastern Pennsylvania calculated an emission rate of between 0.3018-0.41%, a number that accounted for emissions from the production, processing, and transmission of the gas (Peischl et al., 2015). Both of these derived estimates fall below emission rates calculated throughout other basins and are below the 3% threshold required for natural gas to be a smaller climate pollutant in comparison to coal over a 20-year timescale. The low rates in the Marcellus compared to other regions could be the result of a systematic difference within the Marcellus that leads to a more efficient extraction of natural gas. However, while useful as a first-guess estimation, current studies performed in the region are based on relatively small sample sizes (1 aircraft mass balance and 88 individual well measurements). A more thorough analysis of the emission rate in the Marcellus would provide insight into regional differences in CH4 emissions from different shale basins and help improve national estimates of emissions from natural gas.

This study seeks to provide confidence in the emission rate for the northeastern Marcellus by performing the most thorough top-down analysis of the northeastern Marcellus region to date. CH₄ measurements were taken from aircraft observations across 10 flights in northeastern Pennsylvania. A new implementation of modelling CH₄ mole fractions is developed to track complex plume structures associated with different emitters, and an optimal natural gas emission rate is solved for each of the 10 flights. An aircraft mass balance technique is also conducted for 4 of the flights and natural gas emission estimates from this method are compared to those calculated using the modelling technique. Using information on the uncertainty with both methods, a regional emission rate is calculated for the natural gas industry in the northeastern Marcellus region.

137 2 Methods

The objective of this study is to quantify CH₄ emissions coming from unconventional wells and compressor stations, henceforth referred to as upstream natural gas emissions, in the northeastern Marcellus region (defined as the area contain within 41.1-42.2°N 75.2-77.6°W, see Figure 1) through two different top-down methodologies, CH₄ observations from aircraft data are collected for ten (10) individual flights over a three-week period in May 2015. These data are used to solve for the upstream natural gas emission rate using an aircraft mass balance approach. Additionally, a CH₄ emissions inventory for the region is compiled and input into an atmospheric transport model described below. CH₄ concentrations are modelled for each flight, and the upstream natural gas emission rate within the model is optimized to create the best match between aircraft observations and model projected enhancement, providing another estimate for the upstream natural gas emission rate. The sections below detail the regional CH₄ inventory, the aircraft campaign, the transport model, the model optimization technique, and the mass balance approach used in this study.

2.1 Regional Methane Emission Inventory

In this study we characterize emissions from the natural gas industry into five different sectors: emissions from wells, emissions from compressor facilities, emissions from storage facilities, emissions from pipelines, and emissions in the distribution sector.

To estimate CH₄ emissions from the production sector of the natural gas industry, data were first obtained on the location and production rate of each unconventional well from the Pennsylvania Department of Environmental Protection Oil and Gas Reporting website (PADEP, 2016) and the West Virginia Department of Environmental Protection (WVDEP, 2016). To convert the production rate into an emission rate, we need to assume a first-guess as to the expected leakage from wells in the area. A first-guess natural gas emission rate of 0.13% was applied to the production value of each of the 7000+ producing unconventional wells based on the median rate from Omara et al., (2016). The natural gas emission rate was then converted to a CH₄ emission rate by assuming a CH₄ composition in the natural gas of 95% (Peischl et al., 2015).

In addition to unconventional wells, the domain also contains more than 100,000 shallow conventional wells. Annual conventional production rates for the year 2014 were obtained through the PADEP Oil and Gas Reporting website, the WVDEP, and the New York Department of Environmental Conservation (NYDEC, 2016). Despite the large number of wells, the average conventional well in PA produces 1% of the natural gas compared to its unconventional counterpart. However, it is speculated that the older age of these wells and a lack of maintenance and care for them results in a higher emission rate for these wells as a function of their production (Omara et al., 2016). A first-guess natural gas emission rate of 11% was applied to the production values of the conventional wells based on the median emission rate from the wells sampled in Omara et al., (2016). Similar to the unconventional wells, the natural gas emission rate was then converted to a CH₄ emission rate by assuming a CH₄ composition in the natural gas of 95%.

Compressor stations located within the basin are responsible for collecting natural gas from multiple well locations, removing non-CH₄ hydrocarbons and other liquids from the flow, and regulating pressure to keep gas flowing along gathering and transmission pipelines, and can be a potential source for methane emissions. Data for compressor station locations and emissions comes from a dataset used in Marchese et al., (2015). A total of 489 compressor facilities are listed for Pennsylvania, with 87% of the listed facilities also containing location data. Emissions for each compressor station are calculated through two different methodologies. In the simplest case, a flat emission rate of 32.35 kg hr⁻¹ is applied for each station, the mean emission rate of a gathering facility in PA found in Marchese et al., (2015). In the more complex scenario, the same emissions total is used as in the flat rate case, but is distributed among the compressor stations linearly as a function of their energy usage. Wattage between compressors in our dataset can vary greatly, from 10 kW for small compressors to 7000 kW or more at large gathering facilities. Using the wattage as a proxy for emissions allows us to account for the size and throughput of natural gas at each station and assumes larger stations will emit more natural gas compared to smaller stations (Marchese et al., 2015).

Data on locations of underground storage facilities were obtained from the United States Energy Information Administration (EIA, 2015). For each of these locations, a base emission rate of 96.7 kg hr⁻¹ was applied according to the average value emitted by a compressor station associated with an underground storage facility (Zimmerle et al., 2016).

To calculate pipeline emissions, data on pipeline locations needed to be collected. Information on transmission pipelines, which connect gathering compressors to distribution networks, is provided by the Natural Gas Pipelines GIS product purchased from Platts, a private organization which collects and creates various infrastructural layers for the natural gas and oil industry (Platts, 2016). Gathering pipeline data, corresponding to the transfer of gas from wellheads to gathering compressors, is nearly non-existent for PA with the exception of Bradford County, which maps out all gathering pipeline infrastructure within the county border. In PA, information on the location of a gathering pipeline elsewhere is only available where a gathering line crosses a stream or river. To account for gathering pipelines in the remainder of the state, a GIS model was created using Bradford County pipelines map in addition to previously generated pipeline maps of Lycoming County (Langlois et al., 2017) as a typical pattern to simulate connecting pipelines between unconventional wells throughout the state (Figure 2). The resulting pattern follows the valley of the Appalachian Mountains, with larger pipelines crossing through the state to connect the different branches of the network. These pipelines were then multiplied by an emission factor of 0.043 kg per mile of pipe, the factor used for gathering pipeline leaks in the Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2013 (EPA, 2015b).

 CH_4 emissions from natural gas distribution sources, coal mines, and animal/animal waste were provided from Maasakkers et al., (2016), which takes national scale emissions from the EPA's greenhouse gas inventory for the year 2012 and transforms it into a $0.1^{\circ} \times 0.1^{\circ}$ emissions map for the continental U.S. For natural gas distribution emissions, various pipeline data was collected at a state-level and emission factors were accounted for to calculate a total distribution emission for the state. This emissions total was then distributed within the state proportional to the population density. Emission estimates for coal are calculated using information from the Greenhouse Gas Reporting Program (GHGRP) for active mines

and the Abandoned Coal Mine Methane Opportunities Database for abandoned mines (EPA, 2008). State-level emissions missions from enteric fermentation and manure management are provided in the EPA's inventory. These emissions were segregated into higher resolutions using county-level data from the 2012 U.S. Census of Agriculture (USDA, 2012) and land-type mapping.

Finally, the EPA's Greenhouse Gas Reporting Program dataset for the year 2014 was used to capture all other major sources of CH₄ in the region otherwise unaccounted for, the majority of which are emissions from landfills and some industrial sources (EPA, 2015a). Sources within the GHGRP that overlap with natural gas sources already accounted for within our inventory were removed to prevent redundancy.

Although our emissions map used for the model runs did not account for potential CH_4 emissions from wetland sources, a series of wetlands emission scenarios was obtained for the region using data from Bloom, et al., (in review 2017). From this dataset, wetland CH_4 emissions make up only 1% of all regional CH_4 emissions in the most extreme scenario, and thus we assume their impact is negligible to this study.

2.2 Aircraft Campaign

Observations for this project were obtained from a 3-week aircraft campaign during the period of May 14th-June 3rd, 2015and are available for public access (https://doi.org/10.15138/G35K54). The campaign was led by the Global Monitoring
Division (GMD) of the National Oceanic and Atmospheric Administration Earth Systems Research and Laboratory (NOAA
ESRL), in collaboration with the University of Michigan. During this period, the NOAA Twin Otter aircraft flew throughout
the northeast portion of Pennsylvania, providing a total of ten flights across nine days. The aircraft was equipped with a
Cavity Ring-Down Spectroscopic analyser (Picarro G2401-m) measuring CH₄, CO₂, CO, and water vapour mole fractions at
approximately 0.5Hz with a random error of 1 ppb, 0.1 ppm, 4 ppb, and 50 ppm respectively (Karion et al., 2013). GPS
location, horizontal winds, temperature, humidity, and pressure were also recorded at 1Hz1 Hz. The majority of observations
for each flight occurred during the afternoon hours at heights generally-lower than 10001500 m above ground level. Each
flight contains at least one vertical profile within and above the boundary layer, with temperature and water vapour
observations from these profiles used to estimate the atmospheric boundary layer height and ensure that the aircraft sampled
air within the boundary layer throughout the flight. Observations takensuspected of being located above the boundary layer
top are flagged and removed from ealeulationall calculations.

Flight paths, wind speeds, and CH₄ observations for each of the 10 flights can be seen in Figure 3. For six of the ten flights, a box pattern was flown around a large portion of unconventional natural gas wells in northeastern PA. These flights were performed typically on days with a strong, steady wind, with a clearly defined upwind and downwind transect intended for use in an aircraft mass balance calculation. Five of the six box-pattern flights were composed of two loops circling the gas basin, allowing for two separate calculations of the upstream natural gas emission rate for the flight. On the remaining

four flights, raster patterns were performed to help identify spatial complexities of CH₄ emissions within the basin. All ten flights were used in the model optimization calculation of the upstream natural gas emission rate.

2.3 Transport Model

The atmospheric transport model used in this study is the Advanced Weather Research and Forecasting (WRF) model (WRF-ARW, Skamarock et al., 2008) version 3.6.1. The WRF configuration for the model physics used in this research includes the use of: 1) the double-moment scheme (Thompson et al., 2004) for cloud microphysical processes, 2) the Kain-Fritsch scheme (Kain and Fritsch 1990, Kain 2004) for cumulus parameterization on the 9-km grid, 3) the Rapid Radiative Transfer Method for general circulation models (GCMs) (RRTMG, Mlawer et al., 1997, Iacono et al., 2008), 4) the level 2.5 TKE-predicting MYNN planetary boundary layer (PBL) scheme (Nakanishi and Niino 2006), and 5) the Noah 4-layer land-surface model (LSM) that predicts soil temperature and moisture (Chen and Dudhia 2001, Tewari et al., 2004) in addition to sensible and latent heat fluxes between the land surface and atmosphere.

The WRF modelling system used for this study also has four-dimensional data assimilation (FDDA) capabilities to allow meteorological observations to be assimilated into the model (Deng et al., 2009). With WRF FDDA, observations are assimilated through the entire simulation to ensure the optimal model solutions that combine both observation and the dynamic solution, a technique referred to as dynamic analysis. Data assimilation can be accomplished by nudging the model solutions toward gridded analyses based on observations (analysis nudging), or directly toward the individual observations (observation nudging), with a multiscale grid nesting assimilation framework typically using a combination of these two approaches (Deng et al., 2009; Rogers et al., 2013).

The WRF model grid configuration used in this research contains two grids: 9- and 3-km (Figure 4), each with a mesh of 202x202 grid points. The 9-km grid contains the mid-Atlantic region, the entire northeastern United States east of Indiana, parts of Canada, and a large area of the northern Atlantic Ocean. The 3-km grid contains the entire state of Pennsylvania and most of the state of New York. Fifty vertical terrain-following model layers are used, with the centre point of the lowest model layer located at ~10 m above ground level. The thickness of the layers stays nearly constant with height within the lowest 1 kilometre, with 26 model layers below 850 hPa (~1550 m AGL). One-way nesting is used so that information from the coarse domain translates to the fine domain but no information from the fine domain translates to the coarse domain.

The WRF modelling system used for this study also has four-dimensional data assimilation (FDDA) capabilities to allow meteorological observations to be assimilated into the model (Deng et al., 2009). With WRF FDDA, observations are assimilated through the entire simulation to ensure the optimal model solutions that combine both observation and the dynamic solution, a technique referred to as dynamic analysis. Data assimilation can be accomplished by nudging the model solutions toward gridded analyses based on observations (analysis nudging), or directly toward the individual observations (observation nudging), with a multiscale grid-nesting assimilation framework typically using a combination of these two approaches (Deng et al., 2009; Rogers et al., 2013).

FDDA (Deng et al., 2009) was used in this research, with the same strategy as used in Rogers et al., (2013). Both analysis nudging and observation nudging were applied on the 9-km grid, and only observation nudging was applied on the 3-km grid. In addition to assimilating observations and using the North America Regional Reanalysis model as initial conditions, we reinitialize the WRF model every five days, allowing 12 hours of overlapping period in consideration of model spin-up period to prevent model errors from growing over long periods. The observation data types assimilated include standard WMO surface and upper-air observations distributed by the National Weather Service (NWS), available hourly for surface and 12-hourly for upper air, and the Aircraft Communications Addressing and Reporting System (ACARS) commercial aircraft observations, available anywhere in space and time with low-level observations near the major airports.

The WRF model used in this study enables the chemical transport option within the model allowing for the projection of CH₄ concentrations throughout the domain. Surface CH₄ emissions used as input for the model come from our CH₄ emissions inventory and are all contained within the 3-km nested grid. Each source of CH₄ within our inventory is defined with its own tracer (Table 1), allowing for the tracking of each individual source's contribution to the overall projected CH₄ enhancement within the model. For this study, CH₄ is treated as an inert gas. The potential for interaction with the hydroxyl radical (OH), the main sink of CH₄, is neglected. A calculation assuming an above-average OH mole fraction over a rural region of 0.5pptv5 pptv (Stone et al., 2012) and a reaction rate of 6.5×10⁻¹⁵ (Overend et al., 1975) produces a CH₄ sink of 0.5ppb per hour. The duration of a flight can be up to 3 hours, leading to a potential loss of 1.5ppb over the course of a flight. This loss is small but not insignificant, CH₄ plumes associated with natural gas during each flight ranged between 15-70 ppb, and a change of 1.5ppb could theoretically impact observations by as much as 10% of the plume signal. However, this decrease in the CH₄ mole fraction would likely have equal impacts on both the background CH₄ values as well as the enhancement. Because emission calculations are based on the relative difference between the CH₄ background mole fraction and the enhancement downwind, it would take a gradient in the oxidation of OH to impact the results. Considering this relatively low destruction rate, the expected homogeneity of the sink across the region, and the difficulties associated with the simulation of chemical loss processes, we assumed that the CH₄ mass is conserved throughout the afternoon and therefore we ignored the impact of oxidation by OH.

2.4 Model Optimization Technique

2.4.1 Model Optimization Methodology

The objective of the model optimization technique is to solve for a natural gasan emission rate as a percent of natural gas production that creates the best match between modelled CH₄ concentration maps, provided by the transport model, with actual CH₄ mole fraction observations, provided by the aircraft data. The optimization process in this study was originally designed to solve for natural gas emission from unconventional wells and emissions from compressor facilities separately.

Because the flow rate of natural gas being processed was not available for each compressor station, emissions at each facility were originally scaled based on the size of the station. However, when running the transport model using this emissions map, enhancements from the compressor stations produced plume structures nearly identical in shape to enhancements from the unconventional wells due to the similar spatial distributions of these two tracers. Without distinct differences between the enhancement patterns from each tracer, it becomes impossible to distinguish which emissions source must be adjusted to obtain the closest match to the observations. For this reason, emissions from compressor facilities are merged with unconventional well emissions in the optimized emission rate. Though the emission rate solved for in this experiment only uses the locations and production for the unconventional wells, this optimized rate represents emissions from both the wells and compressor facilities and will beare referred to as the modelled upstream natural gas emission rate. Midstream and downstream natural gas processes (such as processing, transmission and distribution of the gas) and emissions from conventional wells are not solved for in this study due to their minimal contribution (less than 5%) to CH₄ emissions in the region encompassed by the aircraft campaign.

Using the transport model WRF-Chem, CH₄ atmospheric enhancements were generated for each flight using different tracers to track different components to the overall CH₄ enhancement (e.g. animal/animal waste, distribution sector, industries). From these concentration fields, the upstream natural gas emission rate was solved for each flight using a three-step model optimization technique. First, a background concentration was determined for each flight and subtracted from the observations to create a set of "observed CH₄ enhancements," using

$$317 \quad X_{EnhO} = X_{Obs} - X_{bg} \qquad , \tag{1}$$

where X_{Obs} is the CH₄ mole fraction observation from the aircraft, X_{bq} is a chosen background value for the flight, and X_{EnhO} is the calculated CH₄ enhancement at each observation. In this study, the background value is defined as the ambient CH₄ mole fraction over the region not accounted for by any of the sources within the model, with each flight having a unique background value. Box-pattern flights containing 2 loops around the basin may have a different background value assigned for each loop. To determine the background mole fraction, we start with the value of the observed mole fraction in the lowest 2nd percentile of all observations within the boundary layer for a given flight or loop. This chosen background value represents the CH₄ mole fraction across the flight path from sources that are outside of our model domain. Because the background value is meant to represent the CH₄ mole fraction outside the model domain whichthat is otherwise unaccounted for in our model, using the observations with the lowest CH₄ mole fraction is not always a sufficient definition for the background. On certain days, CH₄ enhancements from sources within the model domain can form plumes with wide spatial coverage that cover all observations during a flight. For example, during a flight the lowest CH4 observations from the aircraft may be 1850 ppb, but the model simulation during that period indicates that all observations within the flight are being impacted by at minimum a 20 ppb enhancement. In this case, we would set our background value for the flight at 1830 ppb, and say that our 1850 ppb observations from the flight are a combination of an 1830 ppb background in addition to a 20 ppb enhancement from sources within the model. By subtracting off this background value from our observations, we create a set of "observed CH₄ enhancements" which can be directly compared to the model projected enhancements

The next step is to remove enhancements from this set that are not associated with emissions from upstream natural gas using

$$X_{GasO} = X_{EnhO} - X_{OtherM}, (2)$$

where X_{OtherM} is the modelled CH₄ enhancement at each observation from sources unrelated to upstream natural gas processes, and X_{GasO} is the observed observation-derived CH₄ enhancement associated with upstream natural gas emissions for each observation. In this step, each observed CH₄ enhancement has subtracted from it the projected non-natural gas enhancement from the model (i.e. nearest grid point in space) using the corresponding model output time closest to the observation within a 20-minute time interval. This creates a set of observed CH₄ enhancements related only to emissions from upstream gas processes, filtering out potential signals from other CH₄ emitters and providing a set of observed enhancements that can be directly compared to the projected upstream natural gas enhancement within the model. By subtracting these other sources from the observations, we make the assumption that our emissions inventory is accurate for non-natural gas sources and that the transport of these emissions is perfect, both of which are actually uncertain. Because errors exist in both the emissions and transport, it is possible to derivecreate a negative observed observation-derived upstream gas enhancement if model-projected enhancements from other sources are larger than the observed observation-derived enhancement. From the 10 flights, 16% of the observation-derived enhancements are negative, but only 3% are negative by more than 5 ppb. To avoid such situationsolving for unrealistic negative values, these negative observed observation-based upstream gas enhancements are set to 0. Errors associated with this issue and other uncertainties with our inventory are examined further in the results uncertainty analysis section of this paper.

In the final step, the upstream natural gas emission rate within the model is adjusted to create the best match between the modelled upstream gas enhancement and observedobservation-derived upstream gas enhancement using

$$J = \frac{\sum_{k=1}^{n} (X_k^{GasO} - X_{GasO} - C * \frac{X_k^{GasM}}{X_k^2} X_{GasM})}{(3)}$$

where n is the number of observations in the flight, X_{GasO} and X_t^{GasO} and X_t^{GasM} are the observed and modelled enhancement for each observation. In this equation, J is a cost function we are trying to minimize by solving for a scalar multiplier C which, when applied to the modelled natural gas enhancements, creates the smallest sum of the differences between the observed observation-derived upstream gas enhancement and the modelled upstream enhancement. Because the emission rate within the model is linearly proportional to the model enhancements, we can solve for the upstream natural gas emission rate that minimizes the cost function using

$$362 E = 0.13 C (4)$$

where 0.13 was the first guess upstream emission rate (in percent of production) used in the model, and E is the optimized emission rate for the flight as a percentage of the natural gas production at each well. This final value represents an overall emission rate associated with both unconventional wells and compressor stations across the region.

The decision to use a scalar cost function rather than the sum of squares is to account for possible misalignment between any observed CH₄ plume and modelled plumes. There are two potential ways in which misalignment may occur. One possibility is that the modelled wind direction differs from the true wind direction, leading to a plume in the model that is off-centre in relation to the observed plume. The other possibility relates to how the model treats emissions from natural gas as a uniform percent of production. In reality the emissions are more random in nature, and thus the plume may not always develop over the wells with the largest production values. If a cost function is used that minimizes the sum of the squares, any misalignment between the modelled and observed plume will result in the peak of the modelled plume aligning with the height of the tail of the observed plume (Figure 5). Unless the observed plume aligns perfectly with the modelled plume, the optimized emission rate using a sum of squares approach will always bias low. By using a scalar cost function, we solve for an optimized emission rate that results in a plume with the same area under the curve compared to the observed plume (Figure 5). This methodology is not impacted by any misalignment between the modelled vs. observed plumes, preventing the low biases associated with a sum of squares minimization.

2.4.2 Model Optimization Uncertainty Assessment

For each of the ten flights, an uncertainty assessment was performed to obtain a range of likely upstream emission rates for any individual flight. Five different sources of error were considered in this assessment: model wind speed error, model boundary layer height error, CH₄ background error, CH₄ emission inventory error, and model/observation mismatch error. These five sources of error vary substantially from flight to flight depending on conditions, and each can have significant impacts on the total uncertainty (Table 4, 5).

Errors in the modelled wind speed and boundary layer height have impacts on our emission estimates that linearly impact the results. If we assume a constant wind speed, a constant boundary layer height, and no entrainment of air from the top of the boundary layer, we can use the following equation to understand their impacts.

$$\Delta C = \overline{F_0}(\frac{\Delta x}{U * D}) \tag{5}$$

where ΔC is the total CH₄ enhancement of the column of air contained within the boundary layer, $\overline{F_0}$ is the average emission rate over the path the parcel travelled, Δx is the distance the column of air travelled, U is the wind speed and D is the boundary layer height. Using this equation, we can see the linear relationship between the model wind speed, model boundary layer height, and the calculated emission rate. As an example, if wind speeds in the model are biased low, natural gas enhancements projected by the model would increase inversely. To compensate for this effect, the optimized emission rate would decrease proportionally. A similar case can be made for bias in the boundary layer height. Both errors in the wind

speed and boundary layer height have known impacts on the optimized emission rate which can be corrected for, as long as
 the errors of each are known.

To calculate the error in the model wind speed, we assume aircraft observations are truth and use

$$U_e = \frac{\overline{U_{m} - \overline{U_{obs}}}}{\overline{U_{obs}}} \tag{6}$$

where \overline{U}_{obs} is the mean observed wind speed by the aircraft across all points within the boundary layer, \overline{U}_m is the mean modelled wind speed by the model across all points closest in time and space to each observation, and U_e is the wind speed error percentage.

To compute the error in the modelled boundary layer height, the observed boundary layer height for each flight is assumed to be the true boundary layer height and the boundary layer height percentage error, H_S is estimated using:

$$H_e = \frac{H_m - H_{obs}}{H_{obs}} \tag{7}$$

where \overline{H}_{obs} is the average observed boundary layer height across each of the aircraft profiles for a given flight, \overline{H}_m is the model boundary layer height closest in time and space to the location of the observed profiles averaged over all profiles. For both the observation and the model, boundary layer heights were determined by locating height of the potential temperature inversion associated with the top of the boundary layer. On the May 22 flight where a potential temperature inversion could not easily be identified in the observations, changes in water vapour, \overline{CO}_2 and \overline{CH}_4 mixing ratios were used to identify the boundary layer top.

Errors in the model wind speed and boundary layer height are calculated for each of the ten flights. From these errors, a corrected optimized emission rate is calculated for each flight using Eq. (8):

$$E_{new} = \frac{E}{(1+U_e)(1+H_e)} \tag{8}$$

where E is the original emission rate and E_{new} is the corrected optimized upstream natural gas emission rate as a percent of production.

In addition to errors related to wind speed and boundary layer height, we quantify three other sources of error in each flight: errors in the selected CH₄ background value, errors in the CH₄ inventory, and errors associated with the overall model performance (Table 5). Unlike the wind speed and boundary layer errors which have easily computable impacts on the emission estimates, these other three sources of error and their impact on the optimized emission rate are more difficult to quantify.

The background error relates to the value chosen for each flight which represents the ambient CH₄ concentration in the boundary layer unrelated to emission sources within the model. In this study background values ranged from 1897-1923ppb. Though background values should not have high variability during a 2-3 hour mid-afternoon flight, entrainment from the boundary layer top can lead to the mixing in of tropospheric air that has different CH₄ mole fraction values from

those within the boundary layer, resulting in a change in the afternoon background value with time. Furthermore, for days on which all aircraft observations (including those upwind of the unconventional wells) are impacted by various CH₄ plumes predicted within the model, it is difficult to determine the background CH₄ concentration accurately. Additionally, observations corresponding to locations with no modelled enhancement may in fact have been impacted by missing sources in our inventory, highlighting the difficult nature of knowing with certainty where and what the background is for any given flight. Understanding this uncertainty is crucial; any error in subtracting off the background value directly impacts each observation's observed natural gas enhancement. For example, a background value of 1 ppb below the true background for a given flight would add 1 ppb to each observed natural gas enhancement for all observations, creating a high bias with the optimized upstream emission rate. To account for this error, each flight's optimization processes was rerun iterating the background value by ±5 ppb, and the ratio of the percent change in the emission rate compared to the original case was defined as the resulting error in the emission rate due to background uncertainty. This ±5 ppb background error range is an estimate at the range of possible error in the background based on changes observed in the upwind measurements from each of the flights and is meant to be a conservative estimate of the error. The impact this error can have in the emission rate varies depending on the magnitude of the observed downwind enhancements during a flight. A plume containing a CH₄

enhancement of 50 ppb will have a smaller relative error from a 5 ppb change compared to one with an enhancement of only 10 ppb. Thus, days with high wind speeds and a high boundary layer height (and thus enhancements of a smaller magnitude) tend to be affected the most by background errors.

Similar to background errors, errors from the CH₄ emissions inventory are difficult to quantify. In the model optimization technique, we subtract out enhancements from sources unrelated to unconventional natural gas before solving for the upstream gas emission rate. In doing so, we are making the assumption that our emissions inventory for sources unrelated to upstream natural gas processes are accurate. In truth, each emission source in our inventory comes from a different dataset and has its own unique error bounds, many of which are unknown. To simulate the potential errors associated with unknowns in our inventory, we use a Monte Carlo approach and iterate the unconventional emissions optimization approach for each flight 10,000 times, applying a random multiplier between 0-2 for each of the different sources not associated with unconventional natural gas production. The resulting range of optimized natural gas emission rates was fit to a Gaussian distribution and the 2σ emission range was calculated. Despite varying the emissions used in the error analysis by 0 to 200% their original value, their impacts on the optimized natural gas emission rate are minimal on most days due to the northeastern Marcellus region having very few emission sources not related to upstream natural gas processes. Only for the flights on May 24th do we see errors from the inventory contribute significantly to the overall daily

The final source of error attempts to quantify the similarity of the pattern of modelled and observed natural gas enhancements, referred to here as the model performance error. Figure 6 shows an example of two days, one of which the model appears to recreate the observations, and the other of which the model poorly matches the shape of the observed

error, when the coal plume in southwestern PA enters the centre of the study region and has a large role in the upstream

natural gas emission rate calculation for that day (Table 5).

enhancements. Comparing these two simulations with no other information, we hypothesize that one should put more trust in the upstream natural gas emission rate calculated for the flight whose modelled upstream enhancements match structurally compared to the emission rate from the flight whose modelled enhancement bares little semblance to the observed enhancement. The model performance error is designed to account for the trustworthiness of the optimized upstream emission rate based on how well the model simulates a given day. The model performance error is calculated using a modified normalized root mean squared error formula given in Eq. (9):

$$e_{Perf} = \frac{\overline{\sigma}_{\Delta X}}{\Delta X_{QQS}}$$
 (9)

In this equation, $\bar{\sigma}_{\Delta X}$ is the standard deviation of the difference between the modelled and observation-derived upstream natural gas CH₄ enhancement using the optimized emission rate, and ΔX_{gas} is the observed magnitude of enhancement from the major natural gas plume observed in each flight. Here, ΔX_{gas} serves as a normalization factor to account for the varying strength of the enhancement from flight to flight, and ensures that days with increased enhancements due to meteorological conditions or true daily fluctuations in the upstream natural gas emissions do not proportionally impact the performance error percentage. For example, a day with high winds and a deep boundary layer would produce smaller enhancements, leading to a small $\bar{\sigma}_{\Delta X}$ regardless of model performance unless normalized by ΔX_{gas} :

2.5 Aircraft Mass Balance Method and Uncertainty Assessment

An aircraft mass balance calculation was performed for four applicable flights from the aircraft campaign as an alternative method to calculate upstream natural gas emission rates independent of the transport model. The aircraft mass balance approach uses the CH₄ enhancement between a downwind and upwind transect to calculate the total CH₄ flux of the area contained between the two transects. We use the mass balance equation from Karion et al., (2013):

$$479 \mid E = \overline{U}\cos(\overline{\theta}) \int_{-b}^{b} \Delta X \int_{z=0}^{z_{top}} n_{air} dz dx$$
 (510)

where E is the total flux (in mol s⁻¹) coming from the enclosed flight track, \overline{U} is the mean wind speed (in m s⁻¹), $\overline{\theta}$ is the mean angle of the wind perpendicular to the flight track, ΔX is the CH₄ enhancement measured along the downwind flight track from –b to b (expressed as a mole fraction), n_{air} is the molar density of air within the boundary layer (in mol m⁻³), and each of the integrals represents the summing over all air being measured within our transect in both the horizontal (x) and the vertical (z). By simplifying further and using the mean enhancement along each downwind transect as the enhancement and choosing z_{top} to be the top of the boundary layer, we can transform the previous equation into the following:

$$\begin{array}{c|c}
486 & E = 37.3 (L)(D)(\overline{U}) \Delta \overline{X} 3 L D U \Delta \overline{X} \cos(\overline{\theta}) \\
487 & (11)
\end{array}$$

where L is the length of the transect (in meters), D is the depth of the boundary layer (in meters) found using observations from vertical ascents during each flight, $\Delta \bar{X}$ is the mean enhancement across the transect (expressed as a mole fraction), \bar{U} and $\bar{\theta}$ are the mean wind speed (in m s⁻¹) and wind direction relative to the angle of the transect, and 37.3 is the average molar density of dry air within the boundary layer (in mol m⁻³) assuming an average temperature and pressure of 290K and 900hPa.

Flights on May 22nd, May 28th, and May 29th, 2015 were selected for mass balance calculations based on their box shaped flight patterns surrounding a large portion of the northeastern Marcellus gas basin, reasonable steady-state wind conditions during the flight (this eliminated May 14th when a low pressure system was stationed over the region), and a lack of strong CH₄-enhancements originating from outside of the box which could affect both the upwind and downwind transects. Emission rates within the enclosed region were calculated from each of these flights, and emissions not associated with unconventional gas production and gathering sources within the box were subtracted out using information from our inventory. A ratio was taken of the calculated emission rate to the total production in the box, and an upstream natural gas emission rate based on production was obtained. For each of the May 22nd, May 23rd, and May 28th flights, two loops of similar structure and at similar altitudes were performed around the basin, allowing for two individual mass balance calculations on these days. An emission rate is calculated for each of these loops and the two values are averaged to calculate the daily emission rate.

Of the 6 days from the aircraft campaign with a clearly defined upwind and downwind transect, one day (May 14th)-contained a surface high-pressure centre in the middle of the flight resulting in erratic wind patterns, and another day (May 25th) had CH₄ plumes from southwestern PA affecting portions the flight observations. These days were not used for a mass balance, and calculations were performed for the remaining four box-pattern flights (May 22nd, May 23rd, May 28th, May 29th). From this list of remaining flights, three of them contained two loops around a portion of the Marcellus basin. A mass balance was performed on each loop, resulting in a total of 7 mass balance calculations for the region across 4 days. Table 6 summarizes the results from the mass balance flights.

For each flight, a total flux within the box encompassed was calculated using Eq. (11). Using this flux, a natural gas emission rate based on production from within the box was calculated using Eq. (12)

$$E_{0_0} = \frac{E - E_{\text{other}}}{P}$$
 (12)

where E is the total flux from Eq. (11) (in kg hr⁻¹), E_{other} are the emissions enclosed in the box from sources not related to upstream natural gas processes (in kg hr⁻¹), P is the total CH₄ from natural gas being produced within the box (in kg hr⁻¹), and E_{0k} is the resulting natural gas emission rate as a percent of total production within the box.

As an error analysis for the mass balance flight, we look at four potential sources of error (Table 7). One source of uncertainty comes from the observed wind speed used in Eq. (11). For our experiment, we take the mean observed wind speed from the aircraft and assume this value represents the mean wind speed within the entire box during the 2-4 hour period it would take for air to travel from the upwind transect to the downwind transect. To understand the uncertainty and

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biases associated with this assumption, we recreate wind observations along the flight path using values from WRF-Chem, and compare the mean wind speed from the simulated observations to the mean model winds contained within the box integrated throughout the boundary layer during the 3 hour period closest to the flight time. By making this comparison, we are able to understand the representation error associated with treating the wind speed observations from the aircraft as the wind speed within the entire box during the period it would take for air to cross from the upwind transect to the downwind transect. On average, modelled wind speeds following the flight were 7% faster than integrated wind speeds within the box, due to the inability for aircraft observations to account for slower wind speeds closer to the surface. This bias was removed from each day's calculated wind speed. After accounting for the wind speed bias, the average error of the modelled wind speed following the flight path compared to the modelled winds within the box was 3%. This 3% uncertainty was applied to each flight and used as the potential uncertainty in the mean wind speed. Errors in the wind direction were neglected, as each flight used in the mass balance completely surrounded the basin using downwind transects at multiple angles, and thus small errors in the wind angle would result in a negligible net change on the total flux calculated.

Another source of uncertainty is error in the boundary layer height. For each flight, between 2-3 vertical profiles were performed, and the mean height was used in Eq. (11). The standard deviation of different heights from each transect was used as the uncertainty. On May 22^{nd} , a boundary layer height could be interpreted from only one vertical transect. For this day, we assume an uncertainty of ± 200 m ($\pm 9\%$).

Uncertainty in the CH₄ background mole fraction was estimated similar to the boundary layer height. On three of the four flights, two upwind transects were performed. The mean observed CH₄ mole fraction between the two transects was used as the background value for the entire flight, and the standard deviation between the loops was used as the uncertainty. On both the May 23rd and May 28th flights, background differences between the two transects were less than the instrument error of 1 ppb. On these days, we use the instrument error as the background error. On May 22nd, only one upwind transect was usable for the calculation. For this day, we assume a conservative estimate in the uncertainty of the background of ±5 ppb.

Finally, we assess uncertainty in the emissions inventory. After a CH₄ flux is calculated for each loop, emissions from sources contained within the box that are not associated with upstream natural gas processes must be subtracted out to solve for the upstream natural gas emission rate. Any errors associated with our inventory will result in a CH₄ source attribution error. To account for the potentially large uncertainty with the emission sources in our inventory, we vary these non-natural gas emissions by a factor of 2 to test the impact on the solved upstream natural gas emission rate. Because northeastern Pennsylvania contains few sources of CH₄ emissions outside of natural gas production, the impact of this uncertainty is typically less than 20% of the total emissions calculated within the box.

551 3 Results

3.1 Methane Inventory

From the <u>first-guess</u> CH₄ inventory created in this study, a total anthropogenic CH₄ emission rate of 2.76 Tg CH₄ year⁻¹ is projected within our inner model domain (Figure 57) with values for individual source contributions shown in Table 2. This total emissions estimate assumes a leak rate of 0.13% of gas production for unconventional wells, and does not account for emissions from natural gas transmission and storage facilities outside of PA due to a lack of information available from other states. Within the model domain, the area encompassing southwestern PA and northeastern WV stands out as the largest contributor to CH₄ emissions, with emissions from conventional gas, unconventional gas, and coal mines all having significant contributions to the total. In particular, the large emissions from coal make this region unique in comparison to other shales. The EPA's Greenhouse Gas Reporting Program dataset for the year 2014 lists individual coal mines in the southwestern portion of our domain as 8 of the top 10 CH₄ emitting facilities across the entire United States. This large area source of CH₄ can have an impact on CH₄ concentrations hundreds of kilometres downwind and must be taken into account when winds are from the southwest (Figure 68). Examples of this plume and its impacts on the aircraft campaign are discussed in Section 3.2.1.

3.2 Model Optimization Results

3.2.1 Case Studies

From the aircraft campaign, a total of 10 flights across 9 days were used in the model optimization technique. For each one of these flights, CH₄ concentration fields were produced using WRF-Chem, and the emission rate from upstream gas processes was adjusted as outlined in the methods section to find the rate that best matches the total observed CH₄ enhancement. For box flights with two loops completed around the basin, emission rates were calculated for each loop independent from one another and then averaged for the flight. Table 3 provides the general meteorology for the 10 flights.

During each of the observational periods, we use the transport model to project the mole fraction enhancement across the region for each of the different CH₄ tracers (Figure 79). From these projections, we see three common sources of CH₄ which can significantly influence the observed mole fractions in our study region of northeastern PA. The first is emissions from unconventional gas in northeastern PA. Although the first-guess total emissions from upstream production in the Marcellus are small compared to the overall contributions from other sources within the domain, their proximity to the aircraft track results in unconventional gas having the largest contribution to observed enhancements throughout the domain covered by most of the flights, often producing signals downwind of about 20-80 ppb above background levels. The second most influential source of enhancements in our study region comes from various sources of CH₄ emissions located in southwestern PA. Despite being more than 400 km away from our study region, large plumes from coal and other sources in the southwestern corner of the state can contribute enhancements as high as 50 ppb across portions of the flight when winds

are from the southwest, affecting background measurements and masking signals from the unconventional gas. One final, but less influential source of CH_4 enhancement is animal agriculture in southeastern PA. Lancaster County is home to roughly 20% of all cattle in the state, with more than 200,000 cattle and calves as of 2012. A southerly wind can result in a 5-15 ppb enhancement across the flight path due to enteric fermentation and manure management from these cattle. Because of coal, conventional gas, and cattle sources located south of the basin, signals from flights with a southerly component to the wind can be difficult to interpret without modelling the projected plumes associated with these sources. Observations on these days contrast to days with a northerly wind component, where a lack of CH_4 sources north of the study region results in observations with a more clearly defined background and unconventional natural gas enhancement.

For each of the ten flights, variability in the model-observation offset was observed. The first loop of the May 29th flight is the best example of a case where comparisons between the modelled and observed enhancements match closely after optimization. For this flight, a box pattern was flown encompassing a majority of the unconventional wells in northeastern PA, and enhancements were observed along the western and northern transects of the flight. Modelled enhancements from sources unrelated to upstream gas emissions showed a broad CH₄ plume associated mostly with animal agriculture along the western edge of the flight, and a smaller enhancement on the eastern edge associated with two landfills in the Scranton/Wilkes-Barre urban corridor (Figure <u>810</u>). Both of these enhancements are subtracted off from the observations to produce a set of <u>observedobservation-derived</u> enhancements due to <u>upstream</u> natural gas production and gathering facilities. Any enhancements in this new observational dataset are located almost entirely along the northern transect of the flight, directly downwind of the natural gas activity in the region. The <u>observedobservation-derived</u> upstream gas enhancement is then directly compared to the modelled upstream enhancement using its first guess emission rate, and an optimized upstream emission rate of 0.26% of production (i.e. a doubling of the first guess) is calculated by minimizing the difference between the two datasets (Figure <u>911</u>).

The match between observed and modelled CH₄ enhancements on the first loop of the May 29th flight is closer than any other flight in the campaign, with a correlation coefficient of 0.73. The success of the model on this day is likely due to a number of ideal conditions. In general, inconsistencies between the modelled and observed mean wind speeds and boundary layer heights can have a linear bias on the projected enhancements, but for this flight differences between the observed and modelled wind speed and boundary layer height were near 0 for both loops (Figure 10, 1112, 13). Observed wind directions throughout the course of the flight had little spreaddirectional spread and the averaged observed wind direction was only 9° different compared to modelled values, resulting in a transport of the CH₄ plumes that the model was able to match well. Furthermore, the observed mean wind speed was 4.6 m s⁻¹, a moderate wind which allows for a steady transport of any enhancements towards the downwind transect, but not strong enough to dilute their magnitude, resulting in an easily observable enhancement downwind of the basin. Finally, intrusions from sources unrelated to upstream gas were small on this day due to favourable wind conditions, reducing the probability of incorrectly attributing the observed enhancements to the wrong source. Enhancements from upstream natural gas processes were around 30 between 15-40 ppb along our downwind transect. By comparison, enhancements from other sources were lower than 15 ppb along a majority of

the flight, and most of these enhancements were located west of the downwind transect, making them easier to identify and remove without unintentionally impacting enhancements from the natural gas plume. All of these different factors likely contributed to producing a situation where the model was successfully able to match CH₄ observations during the May 29th flight.

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Flights that occurred on days with a southwest wind had a tendency to produce CH₄ observations that were intuitively difficult to interpret due to convolved CH₄ sources in southwestern Pennsylvania. One of these complex observation sets occurred during the late afternoon flight on May 24th, 2015 (Figure 1214). Observations on this day show a CH₄ enhancement-pattern that decreased with latitude, with higher CH₄ mole fractions observed farther south. Given the location of the wells in the middle of the flight path and the WSW wind pattern in the region, this north/south CH₄ gradient is unexpected and counterintuitive compared to where one would expect the enhancements to be based solely on the presence of the gas industry in northeastern PA. However, through modelling each of the many contributors of CH₄ within our inventory, we are able to recreate this latitudinal CH₄ gradient and better understand the observed patterns (Figure 1214). Throughout an 18-hour period leading up to the May 24th flight, winds from the SSW transport emissions from coal in southwestern PA northeastward until they reach the centre of the state, where a westerly wind then shifts the plume across the study region such that it only intersects the southern half of the flight path. Because of both the magnitude of the coal emissions and an accumulation that occurred in the southwestern portion of the state during the previous night, the modelled enhancement from the coal plume is substantial (>20 ppb) as it crosses over the flight path and covers up much of the signal from upstream gas emissions. Nonetheless, the transport model is able to account for these far-reaching sources and attempt to separate out their contribution to the observed enhancements. We are able to-solve for the optimal upstream natural gas emission rate and recreate the May 24th flight observations more accurately than most other flights, with a correlation coefficient of 0.71 between the observations and model values CH₄ values. Although the model successfully recreates the overall observed CH₄ pattern on this flight, attempting to match model vs. observation-derived enhancements specifically from upstream natural gas contributions is much more difficult. Contributions from non-natural natural gas sources are large such that they overwhelm much of the signal from local natural gas sources. After subtracting out non-natural gas sources from the observations, the correlation specifically between modelled and observation-derived upstream natural gas enhancements is only 0.11.

Despite the model's success at recreating observations from the May 24th late-afternoon flight, there is reason to be careful when interpreting results from day with observations influenced by distant sources. In particular, some transport error is unavoidable in atmospheric reanalyses, and the longer the time and distance a plume takes to reach the observations, the more its position and magnitude will be susceptible to these errors. During the early May 24th flight, a small 50 km shift in the location of the coal plume across the study region would change projected enhancements at some observations by as much as 20 ppb. Furthermore, errors in the transport speed could create scenarios where the coal plume either arrives in the study region too early or exits too late, creating a projected enhancement pattern that does not agree with the observations (Figure 1315). Additionally, inaccuracies with the emission estimates of non-unconventional gas sources in the inventory

will impact the magnitude of their CH_4 enhancements, creating additional errors in the optimization process when subtracting out these enhancements from the observations. The early-afternoon May 24^{th} flight and May 25^{th} flight are both examples where influences from CH_4 sources in southwest PA create complex structures in the enhancements, which the model is not able to match as well as the late-afternoon flight on May 24^{th} (Figure 1416). And although observations and modelled enhancements closely match throughout portions of these two flights, a slight shift in the modelled wind direction can lead to vastly differing results due to the large offset small changes in the wind field can have on an emission source hundreds of kilometres away. Thus, results from the flights on May 24^{th} and May 25^{th} should be taken with caution. A deeper analysis of these errors can be found in Section 3.2.2.

Though transport errors from far off sources can have significant impacts on the optimized emission rate, local transport error is handled more effectively in the model through use of the error minimization cost function discussed in the methods section (Eq. 3). Through use of this cost function, we solve for an unstream emission rate that minimizes the area under the curve between the observed and modelled enhancement. This analysis produces a compensation effect that can adjust for misalignment between the observed natural gas CH, plume and the modelled plume, and works best for local sources whose plume structure is similar between the model and observations but location is misaligned. The impact of this effect can be seen best for the flight of May 14th, 2015 (Figure 15). During this day, high pressure was centred just north of the flight track, creating diverging winds that were difficult for the transport model to simulate, and the major plume associated with upstream gas emissions ends up south of a well-defined plume in the observations. Despite this misalignment, the model still simulates the correct magnitude and width of the plume, providing confidence that small errors in the transport of local, well defined structures have only a small impact on the overall emission rate calculation for any . This assumption is corroborated by an analysis of simulated enhancements of an aircraft transecting a plume at an angle of 45° with errors in the projected wind direction varying between ±20°. Such angle would produce errors in the optimized emission rate ranging from ±30% (Figure 16). These errors become smaller as the angle of the aircraft's path with the axis of the major plume approach 90°. Because the downwind transect of most flights is often close to perpendicular with the plume from unstream gas sources, and differences between modelled and observed winds are often less than 20°. sociated with incorrect transport of the CH, plumes would be less

3.2.2 Emission Rates and Uncertainty Assessment

For each of the ten flights, an uncertainty assessment was performed to obtain a range of likely upstream emission rates for any individual flight. Five different sources of error were considered in this assessment: model wind speed error, model boundary layer height error, CH₄ background error, CH₄ emission inventory error, and model/observation mismatch error. These five sources of error vary substantially from flight to flight depending on conditions, and each can have significant imposts on the total presentation.

Errors in the modelled wind direction and boundary layer height have impacts on our emission estimates which have a linear impact on the results. If we assume a constant wind speed, a constant boundary layer height, and no entrainment of air from the top of the boundary layer, we use the following equation to understand these impacts.

$$\Delta C = \overline{F_0} \left(\frac{\Delta r}{H_0} \right) \tag{5}$$

where ΔE is the total CH₄-enhancement of the column of air contained within the boundary layer, $\overline{F_{tt}}$ is the average emission rate over the path the parcel travelled, Δx is the distance the column of air travelled, U is the wind speed and D is the boundary layer height. Using this equation, we can see the linear relationship between the model wind speed, model boundary layer height, and the calculated emission rate. As an example, if wind speeds in the model are biased low, natural gas enhancements projected by the model would increase inversely. To compensate for this effect, the optimized emission rate would decrease proportionally. A similar case can be made for bias in the boundary layer height. Both errors in the wind speed and boundary layer height have known impacts on the optimized emission rate which can be corrected for, so long as the errors of each are known.

To calculate the error in the model wind speed, we assume aircraft observations are truth and use

$$U_{\overline{q}} = \frac{U_{\overline{q}} - U_{\overline{q}}}{U_{\overline{q}}}$$

where \overline{U}_{abc} is the mean observed wind speed by the aircraft across all points within the boundary layer, \overline{U}_{abc} is the mean modelled wind speed by the model across all points closest in time and space to each observation, and U_{ac} is the wind speed error percentage.

To compute the error in the modelled boundary layer height, the observed boundary layer height for each flight is

$$H_{\theta} = \frac{I_{\mathbf{w}} - I_{\mathbf{obs}}}{I_{\mathbf{w}}} \tag{7}$$

where \overline{H}_{am} is the average observed boundary layer height across each of the aircraft profiles for a given flight, \overline{H}_{m} is the model boundary layer height closest in time and space to the location of the observed profiles averaged over all profiles. For both the observation and the model, boundary layer heights were determined by locating height of the potential temperature inversion associated with the top of the boundary layer (Figure 11). In cases where a potential temperature inversion could not easily be identified, changes in water vapour, CO_2 and CH_4 mixing ratios were used to identify the boundary layer top.

Errors in the model wind speed and boundary layer height are calculated for each of the ten flights. From these errors, a corrected optimized emission rate is calculated for each flight using Eq. (8):

$$\frac{F}{E} = \frac{F}{(1+U_{\alpha})(1+U_{\alpha})} \tag{8}$$

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where E is the original emission rate and E_{max} is the corrected optimized upstream natural gas emission rate as a percent of production. Table 4 shows the wind speed and boundary layer height errors for each flight as well as the optimized and corrected natural gas emission rates. On days where model performance was poor in regards to the wind speed and boundary layer height, we can see changes in the corrected emission rate. For most days, this change is less than 20% different than the original optimized emission rate. However, both May 14th and May 25th have corrected emission rates which are around a factor of 2 different from their original value. Whether these corrected emission rates are more accurate than the original optimized rates is debatable. To calculate these alternative emission rates, we must assume that the wind speeds and boundary layer heights from our limited number of observations are the true values in the atmosphere, which may not be the case. Regardless of which rate is more accurate for each flight, the overall 16% high bias in the model wind speed and the ___-12% low bias in the model boundary layer result in compensating errors that cancel out, and the mean emission rates across all flights end up equalsimilar. Thus, any errors associated with these two meteorological variables has a trivial impact on the overall calculated emission rate for the region and the uncorrected emission rates are used for the final mean and uncertainty calculations.

In addition to errors related to wind speed and boundary layer height, we quantify three other sources of error in each flight: errors in the selected CH₄ background value, errors in the CH₄ inventory, and errors associated with the overall model performance (Table 5). Unlike the wind speed and boundary layer errors which have easily computable impacts on the emission estimates, these other three sources of error and their impact on the optimized emission rate are more difficult to quantify.

The background error relates to the value chosen for each flight which represents the ambient CH₄-concentration in the boundary layer unrelated to emission sources within the model. In this study background values ranged from 1897-1923ppb. Though background values should not have high variability during a 2-3 hour mid afternoon flight, entrainment from the boundary layer top can lead to the mixing in of troposphoric air that has different CH₄-mole fraction values from those within the boundary layer, resulting in a change in the afternoon background value with time. Furthermore, for days or which all aircraft observations (including those upwind of the unconventional wells) are impacted by various CH₄-plumer predicted within the model, it is difficult to determine the background CH₄-concentration accurately. Additionally observations corresponding to locations with no modelled enhancement may in fact have been impacted by missing sources in our inventory, highlighting the difficult nature of knowing with certainty where and what the background is for any giver flight. Understanding this uncertainty is crucial; any error in subtracting off the background value directly impacts each observed natural gas enhancement. For example, a background value of 1 ppb below the true background for a given flight would add 1 ppb to each observed natural gas enhancement for all observations, creating a high bias with the optimized upstream emission rate. To account for this error, each flight's optimization processes was rerun iterating the background value by ±5 ppb, and the ratio of the percent change in the emission rate compared to the original case was defined as the resulting error in the emission rate due to background uncertainty. This ±5 ppb background error range is an estimate at the range of possible error in the background based on changes observed in the upwind measurements from each

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 of the flights and is meant to be a conservative estimate of the error. The impact this error can have in the emission rate varies depending on the magnitude of the observed downwind enhancements during a flight. A plume containing a CH, enhancement of 50 ppb will have a smaller relative error from a 5 ppb change compared to one with an enhancement of only 10 ppb. Thus, days with high wind speeds and a high boundary layer height (and thus enhancements of a smaller magnitude) tend to be affected the most by background errors.

Similar to background errors, errors from the CH₄ emissions inventory are difficult to quantify. In the model optimization technique, we subtract out enhancements from sources unrelated to unconventional natural gas before solving for the upstream gas emission rate. In doing so, we are making the assumption that our emissions inventory for sources unrelated to upstream natural gas processes are accurate. In truth, each emission source in our inventory comes from a different dataset and has its own unique error bounds, many of which are unknown. Because of the potential for errors in these emission estimates, we take a conservative approach and iterate the unconventional emissions optimization approach for each flight, varying the emissions from other unsolved for emission sources in the model by a factor of 2 in each direction, thus applying a range of 50 200% to the emissions inventory values to assess its impact on the calculated upstream natural gas emission rate. Despite the extensive range of emissions used in the error analysis, its impact is minimal on most days due to the northeastern Marcellus region having very few emission sources not related to upstream gas processes. Only for the flights on May 24th do we see errors from the inventory contribute significantly to the overall daily error, when the coal plume in southwestern PA enters the centre of the study region and has a large role in the upstream emission rate calculation for that day.

The final source of error accounted for attempts to quantify the similarity of the pattern of modelled and observed natural gas enhancements, referred to here as the model performance error. Figure 17-shows an example of two days, one of which the model appears to recreate the observations, and the other of which the model poorly matches the shape of the observed enhancements. Comparing these two simulations with no other information, we hypothesize that one should put more trust in the upstream natural gas emission rate calculated for the flight whose modelled upstream enhancements match structurally compared to the emission rate from the flight whose modelled enhancement bares little semblance to the observed enhancement. The model performance error is designed to account for the trustworthiness of the optimized upstream emission rate based on how well the model simulates a given day. The model performance error is calculated using Eq. (9):

 $e^{\frac{-\overline{g}_{\pm\pm}}{2}}$

in this equation, $\sigma_{a,c}$ is the standard deviation of the difference between the modelled and observed upstream natural gas CH, inhancement using the optimized emission rate, and ΔX_{gas} is the observed magnitude of enhancement from the major natural gas plume observed in each flight. Here, ΔX_{gas} serves as a normalization factor to account for the varying strength of the enhancements from flight to flight, and ensures that days with increased enhancements due to meteorological conditions.

or true daily fluctuations in the upstream natural gas emissions do not proportionally impact the performance error percentage. For example, a day with high winds and a deep boundary layer would produce smaller enhancements, leading to a small $\bar{\sigma}_{a,k}$ regardless of model performance unless normalized by ΔX_{aux} .

Table 5 summarizes the background error, inventory error, and model performance error, and assumes independence between the three error sources to calculate the total error uncertainty for each flight. The largest error occurred duringuncertainty exists for the May 22nd flight, where an unexplained enhancement along the northern transect led to a poor match between the modelled enhancements and the observed enhancements. This may explain the anomalously high optimized emission rate for that day. Other flights with large error uncertainty are those whichthat occurred on May 24th and May 25th, both days, where enhancements from southwestern PA are believed to be influencing large portions of the observations.

Based on the conservative methodology used to calculate these errorsuncertainties, we assume the total erroruncertainty for each flight represents a 2σ range of possible emission rates and calculate a weighted mean and a 2σ confidence interval for the overall upstream emission rate across the ten flights. From this approach, we find a mean upstream emission rate of 0.36% of production and a 2σ confidence interval from 0.27-0.45% of production.

3.3 Aircraft Mass Balance Results

In addition to the model emission optimization, a simplified aircraft mass balance technique was used to calculated upstream natural gas emission rates for flights with a box pattern, a consistent wind direction within the box, and minimal intrusion of CH₄ enhancements from outside the study region that would affect both the upwind and downwind transects. Of the 6 days with box patterned flights from the aircraft campaign, one day (May 14th) contained a surface low pressure centre in the middle of the flight resulting in erratic wind patterns, and another day (May 25th) had CH₄ plumes from southwestern PA affecting portions the flight observations. These days were not used for a mass balance, and calculations were performed for the remaining box pattern flights (May 22nd, May 23rd, May 28th, May 29th). Of the four remaining flights, three of these flights contained two loops around a portion of the Marcellus basin. A mass balance was performed on each loop, resulting in a total of 7 mass balance calculations for the region across 4 days. Table 6 summarizes the results from the mass balance flights.

For each flight, a total flux within the box encompassed was calculated using Eq. (6). Using this flux, a natural go emission rate based on production from within the box was calculated using Eq. (10)

$$E_{\frac{p}{2q}} = \frac{F - E_{\text{other}}}{P}$$

where E is the total flux from Eq. (6) (in kg hr⁺), E_{other} are the emissions enclosed in the box from sources not related to upstream natural gas processes (in kg hr⁺), P is the total CH_{u} from natural gas being produced within the box (in kg hr⁺), and E_{yy} is the resulting natural gas emission rate as a percent of total production within the box. Calculated emission rates varied extensively between flights used for the mass balance analysis, ranging from 0.11% to 1.04% of natural gas

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production- (Table 6). Comparing emission rates between loops on the same day, we see more consistency in the values. This result is not surprising, as on each of the days with multiple loops, upwind and downwind CH₄ concentrations patterns tended to be similar between loops. Thus, differences in the total emission rate are likely due to either errors specific to each day (such as background variability, errors in meteorology) or real daily variability in the upstream natural gas emission rate.

As an error analysis for the mass balance flight, we look at four potential sources of error (Table 7). One source of uncertainty comes from the observed wind speed used in Eq. (6). For our experiment, we take the mean observed wind speed from the aircraft and assume this value represents the mean wind speed within the entire box during the 2-3 hour period it would take for air to travel from the upwind transect to the downwind transect. To understand the uncertainty and biases associated with this assumption, we recreate wind observations along the flight path using values from WRF Chem, and compare the mean wind speed from the simulated observations to the mean model winds contained within the box integrated throughout the boundary layer during the 3 hour period closest to the flight time. By making this comparison, we are able to understand the representation error associated with treating the wind speed observations from the aircraft as the wind speed within the entire box during the period it would take for air to cross from the upwind transect to the downwind transect. On average, modelled wind speeds following the flight were 7% faster than integrated wind speeds within the box, due to the inability for aircraft observations to account for slower wind speeds closer to the surface. This bias was removed from each day's calculated wind speed. After accounting for the wind speed bias, the average error of the modelled wind speed fellowing the flight path compared to the modelled winds within the box was 3%. This 3% uncertainty was applied to each flight used in the mass balance completely surrounded the basin using downwind transects at multiple angles, and thus small errors in the wind angle would result in a negligible net change on the total flux calculated.

Another source of uncertainty is error in the boundary layer height. For each flight, between 2.3 vertical profiles were performed, and the mean height was used in Eq. (6). The standard deviation of different heights from each transcet was used as the uncertainty. On May 22nd, a boundary layer height could be interpreted from only one vertical transcet. For this day, we assume an uncertainty of ±200m (±9%).

Uncertainty in the CH₄-background mole fraction was estimated similarly to the boundary layer height. On three of the four flights, two upwind transects were performed. The mean observed CH₄ mole fraction between the two transects was used as the background value for the entire flight, and the standard deviation between the loops was used as the uncertainty. On both the May 23rd and May 28th flights, background differences between the two transects were less than the instrument error of 1 ppb. On these days, we use the instrument error as the background error. On May 22rd, only one upwind transect was usable for the calculation. For this day, we assume a conservative estimate in the uncertainty of the background of ±5 ppb.

Finally, we assess uncertainty in the emissions inventory. After a CH₄ flux is calculated for each loop, emissions from sources contained within the box that are not associated with upstream natural gas processes must be subtracted out to solve for the upstream natural gas emission rate. Any errors associated with our inventory will result in a CH₄-source

attribution error. To account for the potentially large uncertainty with the emission sources in our inventory, we vary these non-natural gas emissions by a factor of 2 to test the impact on the solved upstream natural gas emission rate. Because northeastern Pennsylvania contains few sources of CH₄-emissions outside of natural gas production, the impact of this uncertainty is typically less than 20% of the total emissions calculated within the box.

From Table 7, we can see the largest relative errors occurerror with regards to the absolute uncertainty in the emission rate occurs on the May 22nd flight. It is on this day where we have the largest uncertainty in the background value, with observations towards the end of the flight becoming unusable due to a rapid and unexplained decrease in the CH₄ mole fraction with timeof 8 ppb over a 30 minute period (Figure 1817). This day also features the highest boundary layer height and fastest winds of all flights done in this study, reducing the magnitude of the enhancement associated with the natural gas plume and thus amplifying the effects an uncertain background has on the overall uncertainty of the calculated CH₄ flux. Uncertainty across the other three flights is smaller, and results between individual loops on the May 23rd and May 28th flight provide more confidence in the calculated flux for those days.

Using the mean estimated CH_4 emissions and uncertainty for each loop, we calculate a daily mean emission rate and uncertainty for each of the four days. We then solve for an unweighted mean across the four flights to derive our overall emissions estimate from the aircraft mass balance approach, and use the standard error of the flights to estimate the uncertainty. In doing so, we derive a natural gas emission rate from upstream processes of 0.3440% of production, with a 2σ confidence interval from 0.0608-0.6272% of production. Here, we use the arithmetic mean rather than a weighted mean due to the linear relationship between the size of the emission rate and the size of the errors. Because errors associated with ABL height and wind speed have a proportional impact on the calculated emissions within the box, days with a high emissions estimate produce large uncertainties relative to days with a small emission rate. Using a weighted mean approach assigns more weight to the days with low estimated emissions, and produces an overall emission estimate too low and certain to have confidence in $(0.12\pm0.02$ percent of gas production).

4 Discussion

4.1 Upstream Emission Rate

From this study, we estimate with a 2σ confidence interval an emission rate between 0.27-0.45% of gas production using the model optimization method and 0.06-0.62% of gas production using the aircraft mass balance. Figure 1918 provides the emission range estimates from upstream natural gas processes using both the model optimization technique and mass balance technique when applicable. TheseTop-down studies of other basins in the U.S. have all found emission rate estimates as a percentrates greater than 1% of production, and thus the rates calculated for the northeastern Marcellus basin are the lowest observed from top-down measurements of different basins in the U.S., and raiseyet, raising questions as to why thesethe values in the northeastern Marcellus this region appear to be low. One possibility may be related to the well efficiency of the

northeastern Marcellus region compared to other major shale plays (Table 8). In terms of gas production per unconventional well, the Marcellus is the highest of all major basins in the U.S. Furthermore, the gas production per well increases by nearly a factor of two when focusing specifically on Susquehanna and Bradford Counties in northeastern Pennsylvania where the majority of the wells from this study are located (Figure 201). The large difference in production per well between the northeastern Marcellus and other shales may partly explain the low emission rates as a percentage of production. Throughout this study, we normalize natural gas emissions as a percentage of total production under the assumption that higher throughput of natural gas in a system should lead to higher emissions in the system. However, if leaks are more influenced by the number of pneumatic devicescomponents in operation rather than the throughput passing through the devicewells, a high-efficiency production-per-well system such as the unconventional wells in the northeastern Marcellus could end up having a very low emission rate as a percentage of production, but a similar emission rate compared to other basins based on the number of wells, compressors, etc. A thorough bottom-up study of the Marcellus region measuring emissions on a device level could provide an answer to this hypothesis.

Although we calculate a low emission rate for this region, rates calculated for May 22 and May 25 stand out as outliers where emissions fall well-above our uncertainty bounds. It is possible that emissions from natural gas sources were higher on these days compared to others. Releases of natural gas into the atmosphere from short timeframe events such as liquids unloading and venting can add a temporal component to the emission rate. Such events occurring at an increased frequency during the May 22 and May 25 flights could be responsible for the higher emission rates. However, these two days both have issues that could have affected the optimized emission rate. On May 22, we observe a sudden drop in the observed CH_d values that is nearly as large as the main plume on that day, creating concerns about background concentrations. On May 25, a southwesterly wind was present, and while the model showed the coal plume to be west of the flight path, a small shift in the model wind direction would shift the coal plume over the region. For these reasons we are sceptical but not dismissive of the high emission rates found during these two flights.

4.2 Advantages of Combining Observations with Model Output

One of the major advantages of using a chemical transport model to solve for natural gas emission rates compared to a standard mass-balance approach is that the transport model is able to account for the complex and oftentimes non-uniform plume structures originating from sources outside the flight path that can affect observations. When performing a mass balance over a basin, it is assumed that the upwind transect is representative of the air exiting the downwind transect after subtracting out all sources within the box. However, this assumption is only true if winds contained within the flight path are in perfect steady state during the time it take for air to move from the upwind transect to the downwind transect, and that measurements from the downwind transect occurred at a much later time so that the air being measured is the same air measured from the upwind transect. These conditions are not easily achieved for regional scale mass balances due to the long

times needed for the air from the upwind transect to reach the downwind transect. As an example, from the four mass balance flights performed for this study the average time for air to move from the upwind transect to the downwind transect was 4 hours whereas the average time between the aircraft's upwind and downwind measurements was \sim 40 minutes. The aircraft observations can be thought of as a snapshot in time, which can be problematic if large scale plumes from outside the domain are moving through the region and impacting only certain portions of the observations during the flight's short timeframe. By using a transport model for a domain much larger than that of the flight paths, we are able to track these far-reaching plumes and identify situations where the background CH_4 concentrations may be spatially heterogeneous.

The potential usefulness of using a transport model alongside a mass balance calculation can best be demonstrated from observations taken over the Marcellus during a 2013 aircraft campaign (Peischl et. al 2015). During this flight the prevailing winds were from the WSW, and the largest CH₄ enhancements were observed along the western edge of the flight path, upwind of the unconventional wells. Using our transport model, we are able to recreate the day of flight and attempt to use our inventory and explain this feature (Figure 2419). Comparisons between modelled output and observations show a 60 ppb CH₄ enhancement from coal and conventional wells in southwest PA stretching close to the western edge of the aircraft observations, a plume structure similar to the one observed during the May 24th flight from our own study. Though this plume does not initially align with the observed transect with the largest enhancements, we recognize that the coal and gas plume travels for more than 20 hours (a distance of 400 km) from its source before reaching the flight path. If we allow for a 10% error in the transport speed and therefore advance the transport model by an additional two hours past the time in which the aircraft observed these high values, we are able to line up the centre of the plume with the largest observed CH₄ mole fractions along the western edge of the flight. In addition to the 60 ppb enhancement along the centre of the plume, the model projects 20 ppb enhancements along the edges and in front of the plume centre. These smaller enhancements have an influence along different portions of the flight which varies in magnitude, making it difficult to assess a proper background CH₄ value upwind of the wells and potentially masking natural gas enhancements downwind of them. But by using a transport model, we are able to see the potential impact of these far-reaching sources which would otherwise not be considered in a regional mass balance and better understand the complex CH4 plume structures which can occur in a given region under specific wind conditions.

5 Conclusion

 Using the model optimization technique presented in this study, we find a weighted mean natural gas emission rate from unconventional production and gathering facilities of 0.36% of production with a 2 σ confidence interval from 0.27-0.45% of production. This emission rate is supported by four mass balance calculations, which produce a mean of 0.3440% and a 2 σ confidence interval from of 0.0608-0.6272% of production. Applied to all the wells in our study region, this mean rate results in a leakage rate of 20 Mg CH₄ hr⁻¹, for the year 2015. The emission rate found in this top-down study quantified as a percent of production is significantly lower than rates found using top-down methodology at any other basin, and indicates

the presence of some fundamental difference in the northeastern Marcellus gas industry that is resulting in more efficient extraction and processing of the natural gas.

The ten flights that took place in this study reveal large regional variations in the CH₄ enhancement patterns depending on the prevailing wind direction. On days with a northwest wind, observed enhancements come primarily from the natural gas industrysources, and a small plume associated with it can be seen on the downwind leg of each flight with few enhancements upwind of the wells. Flights whichtjat took place with winds conditions predominantly from the southwest were more difficult to interpret. Plumes associated with coal and other potential sources of CH₄ in the southwestern Pennsylvania create complex enhancement patterns affecting both the upwind and downwind portions of the flight, making both the background CH₄ mole fraction and enhancements from the gas industry difficult to interpret. The stark difference between observations whichthat occurred with a northwest wind compared to a southwest wind illustrates the importance of having multiple flights across days with various wind conditions to better understand the major influences on CH₄ concentrations throughout a region. The regional influences in Pennsylvania also demonstrate the utility of deriving an emissions inventory that provides input data to drive a transport model, allowing one to forecast CH₄ mole fractions on difficult days and better understand the daily uncertainties associated with heterogeneous background conditions.

Though this study presented observations from ten flights over a three-week period, it is not able to account for the potential of long term temporal variability in the emission rates. In May 2015 when the flights took place, the entire Marcellus basin was nearing peak production and active drilling and hydraulic fracturing was still ongoing in the region. By mid-2016, the rate of drilling of new wells in the northeast Marcellus had decreased and natural gas production had begun to decline in the area. A snapshot of the emission rate during one month of a basin in its peak production is insufficient to characterize emissions from an area that is likely to be producing and transporting gas at various intensities for decades. We need to quantify the long-term climatological impacts of gas production. Future work examining the temporal variability of CH₄ emissions within natural gas basins would complement short-term, high-intensity studies such as this one, and aid with understanding how well the calculated emission rates represent the gas basin over the course of time.

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Table 1: List of tracers used in the transport model.

Tracer #	Name	Description of source
1	Unconventional Wells	Emissions from unconventional wells.
2	Storage Facilities	Emissions from compressors associated with natural gas storage.
3	Pipelines	Emissions from gathering and transmission pipelines
4	Distribution	Emissions from the distribution sector of the natural gas industry.
5	Conventional Wells	Emissions from conventional wells.
6	Landfills/Other	Emissions from landfills and uncharacterized industrial sources.
7	Coal	Emissions from active and abandoned coal mining.
8	Animals/Waste	Emissions from enteric fermentation and manure management
9	Production Compressors (HP)	Emissions from compressor stations characterized as "production". Emissions scaled linearly with wattage.
10	Gathering Compressors (HP)	Emissions from compressor stations characterized as "gathering". Emissions scaled linearly with wattage.
11	Other Compressors (HP)	Emissions from all other compressor stations. Emissions scaled linearly with wattage.
12	Production Compressors (C)	Emissions from compressor stations characterized as "production". Emissions constant among compressors.
13	Gathering Compressors (C)	Emissions from compressor stations characterized as "gathering". Emissions constant among compressors.
14	Other Compressors (C)	Emissions from all other compressor stations. Emissions constant among compressors.

Table 2: Annual emission rate totals from anthropogenic sources within the innermost model domain based on values from the inventory within this study

Source	Total Emission Rate (Gg CH ₄ year ⁻¹)
Unconventional Wells	125
Conventional Wells	607
Gathering Compressor Facilities	118
Storage Facilities	69
Gathering/Transmission Pipelines	8
Natural Gas Distribution	213
Underground, Surface, and Abandoned Coal Mines	831
Enteric Fermentation/Manure Management	371
Landfills	420
Total	2762

Table 3: Meteorological statistics from the May 2015 flight campaign.

Day	Flight	# of	# of	ABL	Mean Observed	Mean	Model
	Pattern	Loops	Vertical	Depth	Wind Speed	Observed	Background
			Profiles	(m)	(m/s)	Wind	Value (ppm)
						Direction	
May 14	Box	1	2	1300	2.9	30°	1.908
May 21	Raster	N/A	2	1300	3.9	231°	1.905
May 22	Box	2	2	2300	10.1	300°	1.910
May 23	Box	2	2	1400	4.4	276°	1.906
May 24 ¹	Other	N/A	2	1500	4.4	270°	1.923
May 24 ²	Raster	N/A	2	2050	4.8	272°	1.907
May 25	Box	1	2	1800	9.0	217°	1.920
May 28	Box	2	3	1400	7.1	322°	1.897
May 29	Box	2	2	1000	4.6	195°	1.899
June 3	Raster	N/A	1	1250	2.7	149°	1.898

Table 4: Optimized natural gas emission rates for each flight as well as corrected emission rates adjusting for errors in the model wind speed and boundary layer height. For wind speed and boundary layer height error, a negative value represents a model value less than the observations.

Day	Optimized NG	Wind	Boundary	Corrected NG
	Emission Rate (%	Speed	Layer Height	Emission Rate (% of
	of production)	Error (6)	Error (7)	production)
May 14	0.37	-31%	-33%	0. 17 <u>80</u>
May 21	0.53	3%	39%	0. 76 <u>37</u>
May 22	1.15	37%	-18%	1. 30 <u>02</u>
May 23	0.45	34%	-9%	0. 55 <u>37</u>
May 24	0.68	48%	-21%	0. 80 <u>58</u>
May 24	0.36	48%	-21%	0. 42 <u>30</u>
May 25	0.99	3%	-43%	0.58 1.69
May 28	0.33	-4%	-8%	0. 29 <u>37</u>
May 29	0.35	4%	1%	0. 37 <u>33</u>
June 3	0.26	19%	-8%	0. 29 24
Average	0.55	16%	-12%	0. 55 <u>61</u>

Table 5: Emission rates and potential errors associated with the model optimization technique. <u>r-values represent the correlation between the model and observation-derived upstream natural gas enhancements.</u>

Day	Optimized	<u>r-value</u>	Background	Non-Upstream	Model	Total	2σ
	Upstream	Model	Error	Gas Inventory	Performance	Error	Confidence
	Emission	vs Obs		Error	Error		Interval
	Rate (% of	<u>NG</u>					(% of
	production)	Sources					Production)
May 14	0.37	0.20	±24%	± 20 19%	±17%	± 36 35%	±0.13
May 21	0.53	0.31	±24%	± 17 13%	±30%	±4241%	±0.22
May 22	1.15	0.47	±38%	± 6 5%	±37%	±53%	±0.61
May 23	0.45	0.10	±39%	±13%	±42%	±59%	±0.26
May 24 ¹	0.68	0.31	±24%	± 54 <u>81</u> %	±17%	± 61 <u>86</u> %	±0.4258
May 24 ²	0.36	0.11	±51%	± 78 150%	±31%	± 98 161%	±0. 35 57
May 25	0.99	0.43	±29%	± 19 15%	±30%	±4644%	±0. 45 44
May 28	0.33	0.33	±76%	± 36 12%	±20%	± 86 79%	±0. 29 26
May 29	0.35	<u>0.58</u>	±24%	± 9 11%	±19%	± 32 33%	±0. 11 12
June 3	0.26	0.37	±31%	± 10 12%	±24%	±4041%	±0.11

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Table 6: Emission rates from mass balance calculations on applicable days, with emission ranges associated with a ± 5 ppb error in the background value.

Flight	CH ₄	Mass	Non-	Calculated Upstream	2σ Confidence
	Production	Balance	Upstream	Emission Rate (% of	Interval (% of
	within box	CH ₄ Flux	CH_4	production)	Production)
	(Gg hr ⁻¹)	(kg hr ⁻¹)	Emissions		
			(kg hr ⁻¹)		
May 22 ₁	4.96	53800	2250	1.04	±1.09
May 22 ₂	4.96	27400	2250	0.51	±1.08
May 23 ₁	4.05	5600	934	0.11	±0.07
May 23 ₂	4.05	5500	934	0.11	±0.07
May 28 ₁	3.73	7100	706	0.17	±0.11
May 28 ₂	3.73	6000	843	0.14	±0.10
May 29 ₁	4.63	27900	1622	0.57	±0.30

Table 7: Relative error associated with the different sources of uncertainty in the aircraft mass balance.

Flight	Wind	Background	ABL Error	Inventory	Total	Upstream Emission
	Speed	Error		Error	Error (1σ)	Rate (% of
	Error				. ,	Production) w/ 2σ
						Confidence Interval
May 22 ₁	±3%	±56%	±9%	±5%	±57%	1.04 ±1.09
May 22 ₂	±3%	±121%	±9%	±8%	±121%	0.51 ±1.08
May 23 ₁	±3%	±24%	±7%	±20%	±32%	0.11 ±0.07
May 23 ₂	±3%	±26%	±7%	±21%	±34%	0.11 ±0.07
May 28 ₁	±3%	±31%	±7%	±11%	±34%	0.17 ±0.11
May 28 ₂	±3%	±33%	±7%	±16%	±38%	0.14 ±0.10
May 29 ₁	±3%	±28%	±20%	±8%	±36%	0.57 ±0.30

Table 8: Production statistics from mid-2014 for various shales across the United States (Hughes 2014).

	Barnett	Fayetteville	Haynesville	Marcellus	Bradford/ Susquehanna County, PA
# of Producing Wells	16100	4500	3100	7000	1558
Total Production (Bcf day ⁻¹)	5.0	2.8	4.5	12	5.01
Production per well (MMcf day ⁻¹)	0.31	0.56	1.25	1.71	3.22

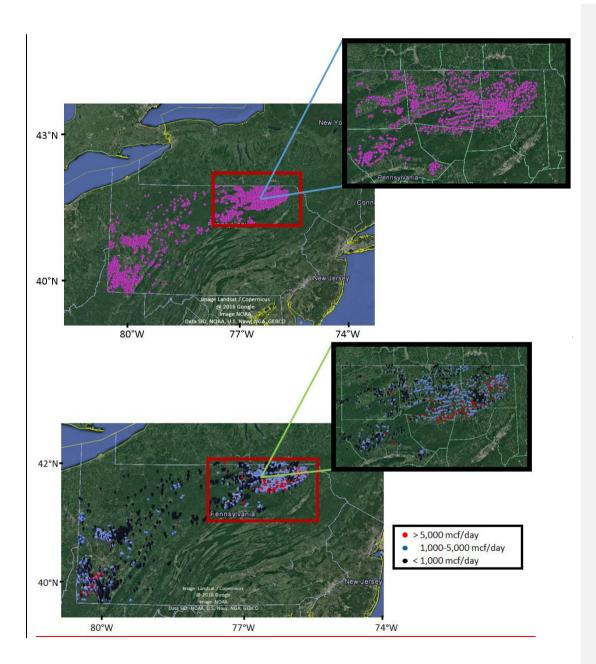
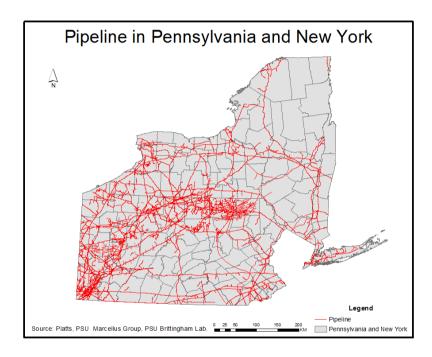


Figure 1: A map of the unconventional wells in Pennsylvania dotted in purple. Production values of wells for May 2015 are indicated by the marker colour. Red rectangle and zoom-in show the region of focus for this study, 41.1-42.2°N 75.2-77.6°W.



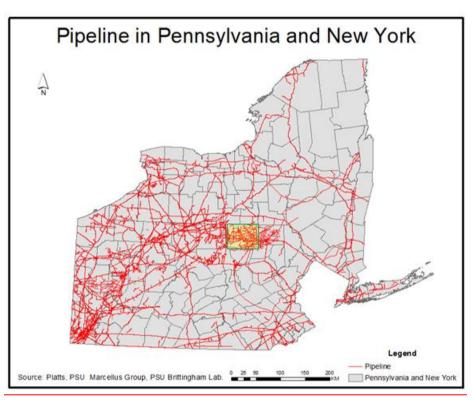


Figure 2: A map of transmission and gathering pipelines for the state of PA and NY. Transmission pipelines are provided by Platts Natural Gas Pipelines product. Gathering pipelines associated with unconventional wells in PA are extrapolated using information on existing gathering pipelines provided by Bradford County, PA_{τ} (highlighted in yellow).

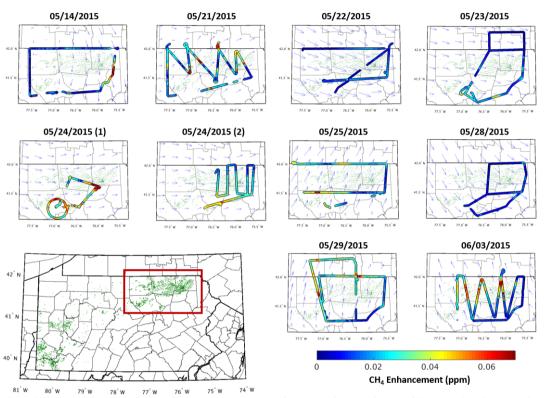


Figure 3: Observed CH_4 enhancements within the boundary layer from each of the 10 afternoon flights used in this study, with green dots showing the location of unconventional wells in PA and blue arrows showing the modelled wind direction during the time of the flight. CH_4 enhancements are calculated by taking the observed CH_4 mole fraction values and subtracting off the flight's background CH_4 value shown in Table 3.

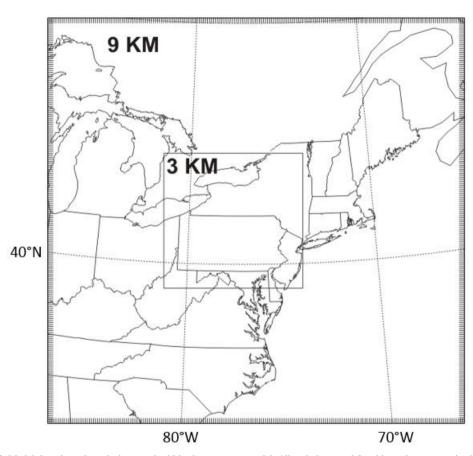


Figure 4: Model domain and resolutions used within the transport model. All emissions used for this study are contained within the $3 \mathrm{\ km}$ resolution domain.

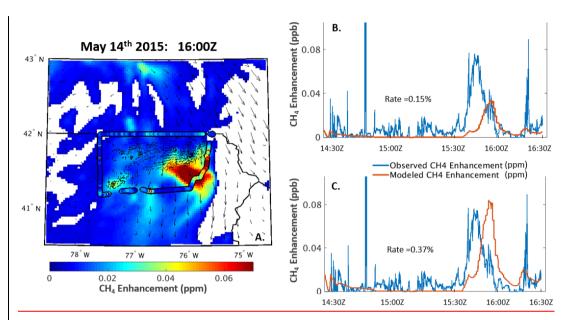


Figure 5: (a.)

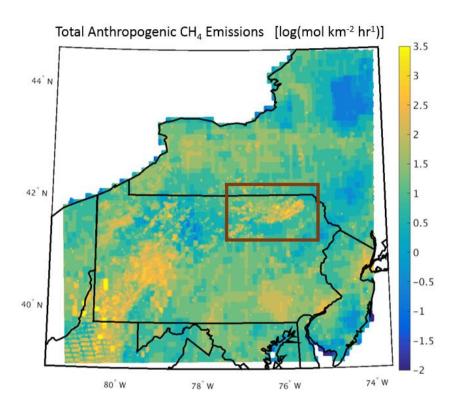


Figure 5 Observed vs model projected CH₁ enhancements during the May 14th, 2015 at 16Z. (b.) Comparison of observed natural gas enhancement to modelled natural gas enhancement along flight path, with upstream emission rate optimized by minimizing the absolute error between the datasets. (c.) Same as previous, but optimized by minimizing the sum of the error between the datasets.

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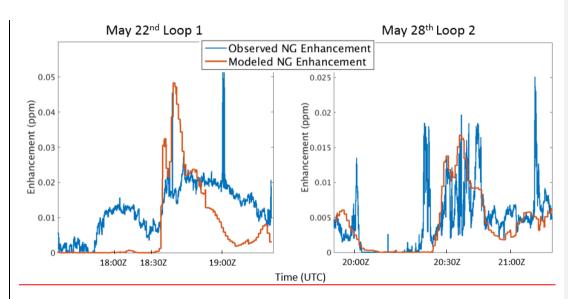


Figure 6: Comparison of observed natural gas enhancement to modelled natural gas enhancement for segments along the (left) May 22nd flight and (right) May 28th flight. A distinct lack of representativeness of the observations in the modelled enhancement can be seen in the May 22nd flight compared to the May 28th flight.

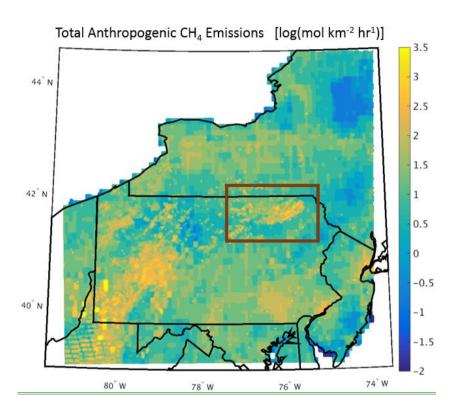


Figure 7: A log scale contour of the anthropogenic CH_4 emissions inventory from this study used within the transport model. The red rectangle surrounds the study region where the aircraft campaign took place.

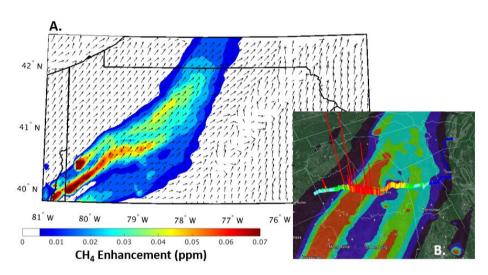


Figure $\frac{68}{1}$: (a.) Model projected CH₄ enhancement at the surface associated with underground, surface. and abandoned coal mines on May 27^{th} , 2015 at 19Z, with the shaded regions showing the CH₄ enhancement and the arrows representing the wind direction. (b.) Projected enhancement from a mapped over measured CH₄ enhancement from a driving campaign. The height and colour of the bars represents the scale of the CH₄ enhancement.

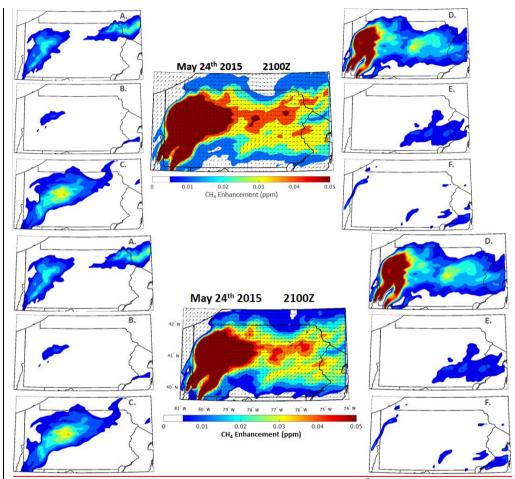


Figure 79: Projected CH₄ enhancements during the late afternoon flight of May 24th, 2015 at 2100Z, 700m above ground level from (A) upstream unconventional gas processes (B) downstream unconventional gas processes (C) conventional production (D) coal mines (E) animal emissions and (F) landfills and other sources within the EPA GHG Inventory Report. The centre figure is a map of the combined enhancement from sources A-F.

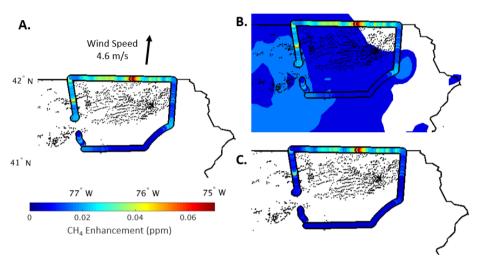


Figure $8\underline{10}$: (a.) Observed CH₄ enhancements from within the boundary layer during the first loop of the May 29^{th} aircraft campaign. (b.) Aircraft observations laid overtop modelled CH₄ concentrations at 700 m from sources unrelated to emissions from upstream gas production. (c.) Observed CH₄ enhancements from the May 29^{th} flight after subtracting off modelled sources in b. The new set of observations represent the observation-derived upstream gas enhancement during the flight.

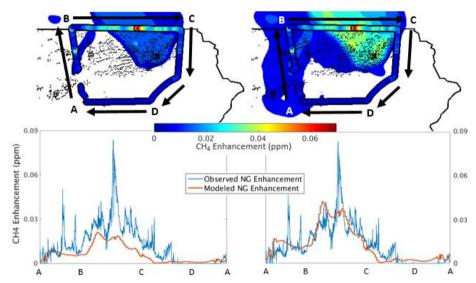


Figure 911: (top-left) Observed enhancement from unconventional natural gas production overtop projected upstream natural gas enhancements at 700 m from the first loop of the May 29th flight, using an upstream gas emission rate of 0.13% of production. (bottom-left) Direct comparison of the observed natural gas enhancement vs. the modelled enhancement following the path from A-D using an unconventional emission rate of 0.13%. (top-right, bottom-right). Same as left figures, except using the optimized upstream emission rate of 0.26%

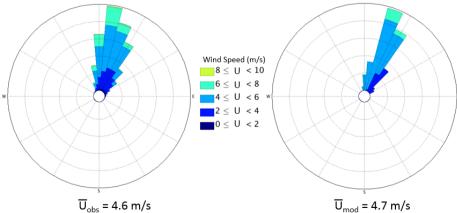


Figure <u>4012</u>: Wind rose of aircraft observations (left) within the boundary from the first loop of the May 29th flight compared to modelled winds following the flight path (right).

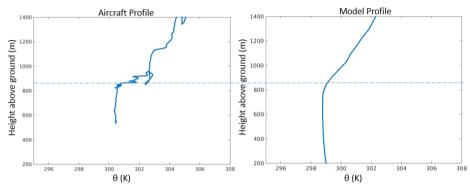


Figure 4413: (left) Observed potential temperature profile with height from the first aircraft spiral on the May 29th flight at 17Z. (right) Modelled potential temperature at the location and time at which the aircraft spiral occurred. In both cases, an inversion in the potential temperature profile begins to occur around 850m.

May 24th 2015: Late-Afternoon Flight

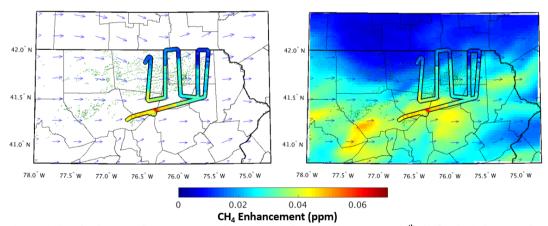


Figure 1214: (left) Observed CH₄ enhancement from the late-afternoon flight on May 24th, 2015. (right) Observed CH₄ enhancement compared to the model projected CH₄ enhancement from the sum of all sources in the region. The colour scale of observed and projected enhancements is scaled 1:1, with matching colours indicating matching values. Modelled wind vectors and CH₄ concentrations are from 700 m model height level.

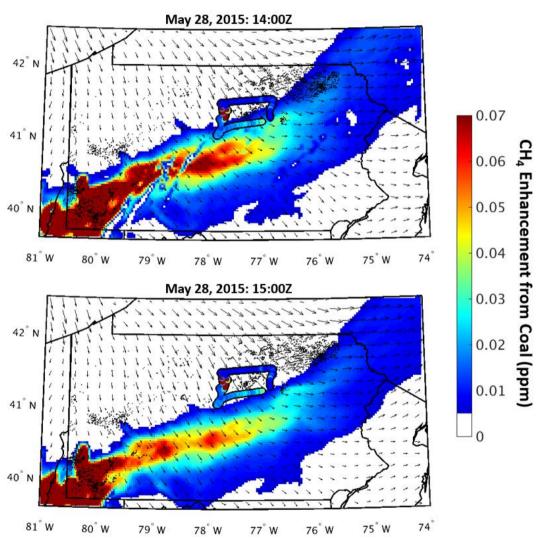


Figure 1315: Observed CH4 enhancements from an early flight on May 28th, 2015 compared to projected CH4 enhancements from coal emissions modelled at (top) 14:00Z and (bottom) 15:00Z. The one hour time difference results in vastly different projected enhancements across the southern portion of observations. Modelled wind vectors and CH₄ concentrations are from the 700 m model height level.

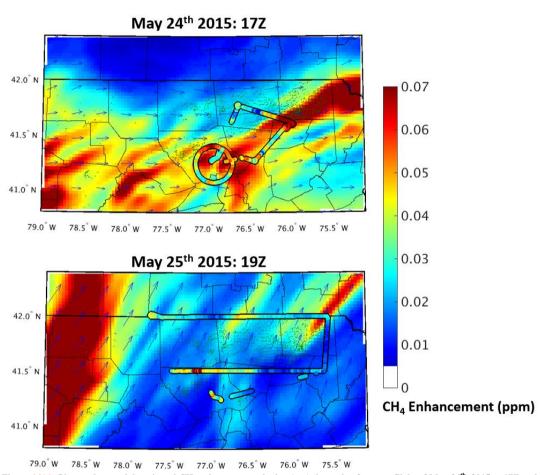


Figure 4416: Observed vs model projected CH_4 enhancements during (top) the early afternoon flight of May 24^{th} , 2015 at 17Z and (bottom) the flight of May 25^{th} , 2016 at 19Z. Modelled wind vectors and CH_4 concentrations are from 700 m model height level.

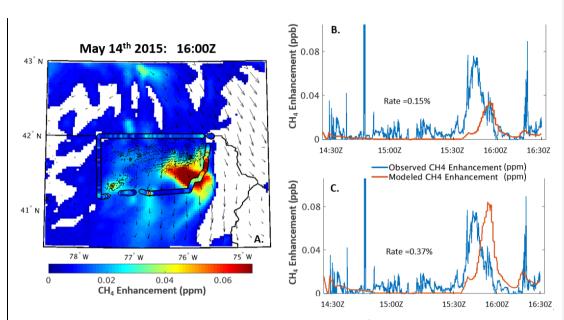


Figure 15: (a.) Observed vs model projected CH₄ enhancements during the May 14th, 2015 at 16Z. (b.) Comparison of observed natural gas enhancement along flight path, with upstream emission rate optimized by minimizing the absolute error between the datasets. (c.) Same as previous, but optimized by minimizing the sum of the error between the datasets.

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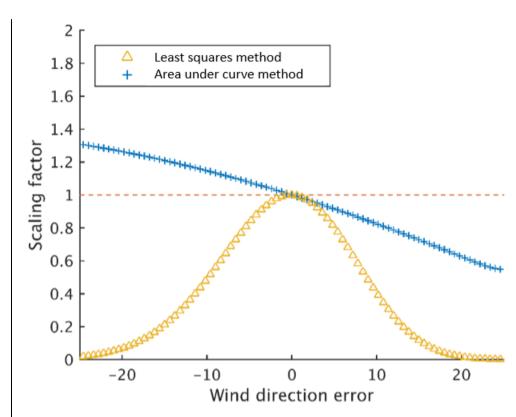
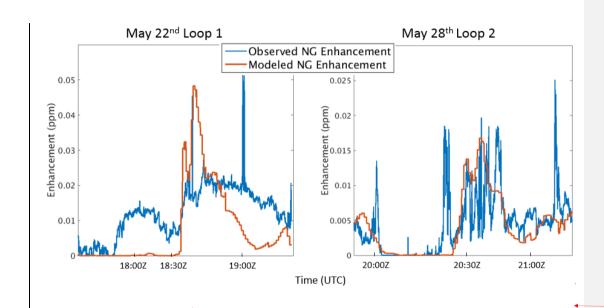


Figure 16: A demonstration of the effect of model transport error on the optimized emission rate using pseudo-observations transecting a modelled plume at a 45° angle. The scaling factor represents the change in the optimized emission rate compared to the true emission rate, with 1 representing a perfect rate.



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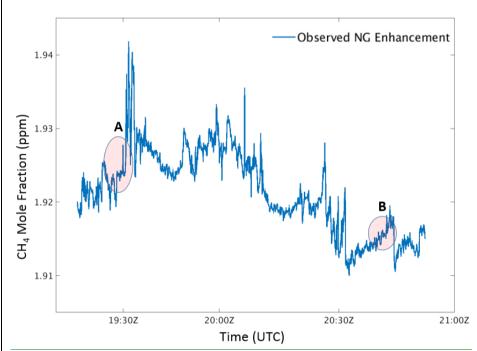


Figure 17: Comparison of observed natural gas enhancement to modelled natural gas enhancement for segments along the (left) May 22nd flight and (right) May 28th flight. A distinct lack of representativeness of the observations in the modelled enhancement can be seen in the May 22nd flight compared to the May 28th flight.

Figure 17

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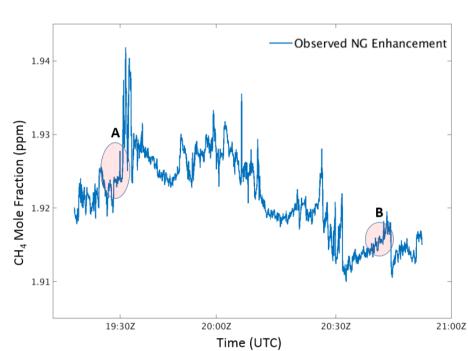
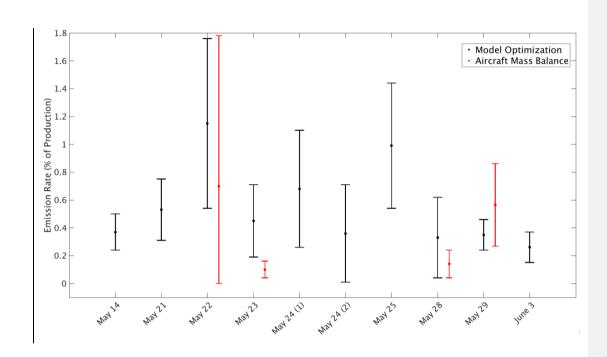


Figure 18: Time series of CH_4 mole fractions from the second loop of the May 22^{nd} flight. Observations at the shaded areas below A and B were taken at similar locations in space, showing the change in the background mole fraction across time.



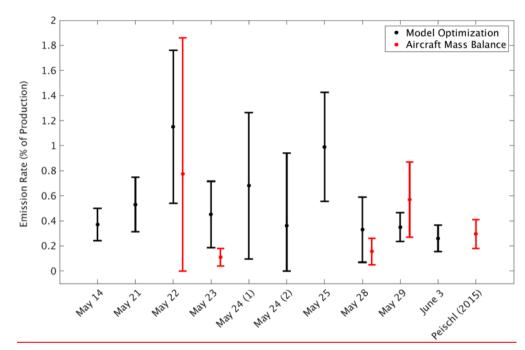


Figure $\frac{1918}{2}$: Calculated upstream natural gas emission rates using (black) model optimization technique and (red) aircraft mass balance technique. Error bars represent the 2σ confidence interval for each flight. Mass balance performed in Peischl et al (2015) included for comparison.

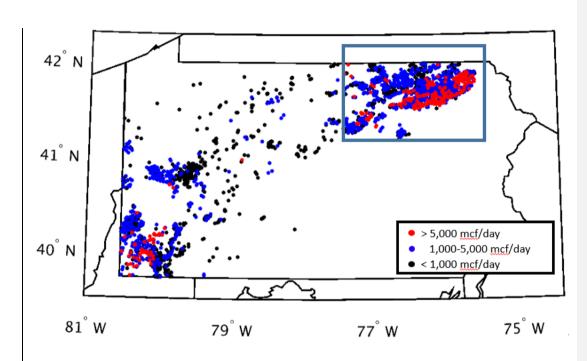


Figure 20: Well locations and daily production of unconventional wells in PA for May 2015. Boxed region is study region where flights took place.

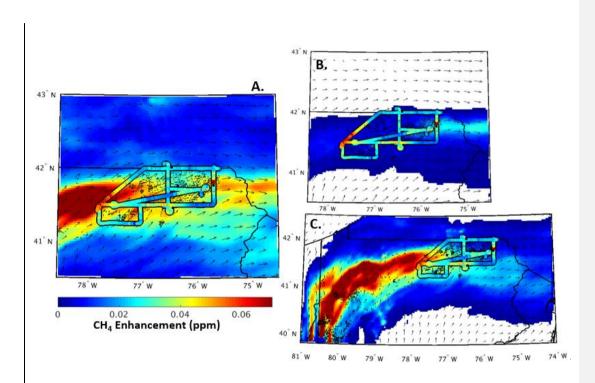


Figure 21

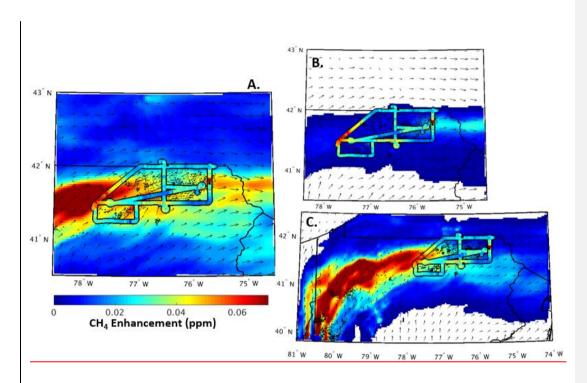


Figure 19: Observations vs modelled enhancements of the flight from Peischl et. al (2015) for July 6th, 2013. (a.) Observed enhancements from the flight over model projected enhancements from all sources at 21Z. (b.) Projected enhancement from upstream gas processes using a 0.4% emission rate. (c.) Projected enhancement from coal sources in southwestern PA. Modelled wind vectors and CH₄ concentrations are from 700 m model height level.