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Title: Methane emissions from oil and gas platforms in the North Sea

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Dear Editor,

We thank the referees and editor for their comments, especially the additional data supplied by Reviewer 1. As suggested, we have amended the manuscript to address the reviewers' comments and suggestions.

Please find our detailed responses below.

Yours sincerely,

Stuart Riddick (corresponding author)

Stuart N. Riddick, Denise L. Mauzerall, Michael Celia, Neil R. P. Harris, Grant Allen, Joseph Pitt, John Staunton-Sykes, Grant L. Forster, Mary Kang, David Lowry, Euan G. Nisbet, and Alistair J. Manning

Anonymous Referee #1

General comment

Reviewer Comment:

I have reservations about: 1) the applicability of the Gaussian plume model to the conditions, though these reservations could well be ameliorated with additional information and author responses. 2) The relevance of the observations made at the coast 3) the quality of the results of the Keeling plot analysis 4) the uncertainty analysis of the Gaussian plume model

I may be missing it, but I can't find public access to much of the data in this paper. Best practices for reproducibility would have all data publicly accessible with access instructions given in the paper. I would appreciate the opportunity to inspect and analyze the data before making the recommendation to publish. There are a few places in the paper with insufficiently detailed information to fully understand the study - for example, what are the coordinates of the platforms and on what day was what platform observed? How much did winds vary over the course of the ship transects?

Author's response:

In the responses below we address the reviewer's reservations regarding the suitability of the Gaussian plume model and the uncertainties. We also acknowledge the reviewer's concerns regarding the uncertainty analysis and have enhanced the content of this section to indicate the shortcomings of the measurement methodologies. To address the issue of data transparency we have introduced all data required to reproduce the emission estimates in Supplementary Materials Sections 1 and 2 and included latitudes and longitudes of the measured platforms.

With regard to the relevance of the observations made at the coast and the quality of the results of the Keeling plot analysis we have decided to remove these sections completely and make the focus of this manuscript solely on offshore measurements.

Critique #1

The use of a Gaussian plume model requires careful consideration of the assumptions that go into such a model. Namely, the model assumes a homogenous, steady state flow with a steady point source.

I think that, for the most part, the conditions in this study satisfy those assumptions, but I do have the following reservations.

The Gaussian plume model employed by the authors assumes an infinitely high boundary layer and homogenous mixing throughout the boundary layer which is to say that they include a reflection term at the surface, no reflection term at the top of the boundary layer, and a uniform vertical mixing. This is a marine environment in a cool climate during the summer, and so a marine layer is likely. This would come with a very low boundary layer height and temperature inversion near the ocean surface. The emission heights are 50-70m and the inlet height is 2.5m. The assumptions of homogenous vertical mixing and no reflection off the top of the boundary layer are at risk.

I had code on hand to extract and plot meteorological fields from the GFS global forecast model archives (raw data obtained from <https://ready.arl.noaa.gov/archives.php>). I extracted boundary layer heights and winds at 1300UTC on the days of the campaign and plotted them below. Boundary layer height capped at 1500m in the plot for visibility. These data carry the caveat that GFS archive forecast data has error – particularly in the boundary layer height. I am happy to share these data/plots with the authors for their own use if they wish.

The paper includes a plot of the studied platforms, but no quantitative description of the locations (e.g., latitude and longitude, and which day each was measured). However, it does appear that the boundary layer height in the vicinity of the platforms may have been quite low, depending on when each was observed.

Author's response

We thank the reviewer for highlighting issues with the Gaussian plume model and in particular for assisting our analysis by sharing data. We greatly appreciate these contributions and are confident that our additions have improved the manuscript. To address the reviewers concerns we have included the assumptions we have made when using the Gaussian plume model.

To specifically address the concerns over the plume's reflectance at the top of the boundary layer we have modified the Gaussian Plume model that we use to include terms that account for reflectance at the top of the boundary layer. The height of the boundary layer is taken from the Global Forecast System's global forecast model archives provided by the reviewer. These data are presented in the full data table in Supplementary Material Section 1, where all of the data used to calculate emissions from each platform can be found.

To address the issue of homogeneous mixing the shortcomings of the Gaussian plume model have been added to new sections: 2.2.5 Uncertainties; and 3.5 Uncertainties/shortcomings of Gaussian Plume modelling.

Change to manuscript:

P5 L8

“the height of the source (h_s , m), **the height of the boundary layer (h , m)** and the stability of the air (CERC, 2017; Hunt, 1982; Hunt et al., 1988). The standard deviation of the lateral (σ_y , m) and vertical (σ_z , m) mixing ratio distributions are calculated from the Pasquill Gifford stability class (PGSC) of the air (Pasquill, 1962; Busse and Zimmerman, 1973; US EPA, 1995). **Even though this modelling method is relatively simple, offshore emissions estimates using the same parameterization of σ_y and σ_z were made by Blackall et al. (2008) and were in good agreement ($R^2 = 0.85$) with emissions calculated from a concurrent tracer release experiment. Alternative offshore parameterisations for σ_y and σ_z exist and are used in the EPA recommended Offshore and Coastal Dispersion Model (Hanna et al, 1985). However, these algorithms require further data on the micrometeorology which are not available and were therefore not used as they introduce additional unquantifiable uncertainty.**

$$X(x, y, z) = \frac{Q}{2\pi u \sigma_y \sigma_z} e^{\frac{-y^2}{2\sigma_y^2}} \left(e^{\frac{-(z-h_s)^2}{2\sigma_z^2}} + e^{\frac{-(z+h_s)^2}{2\sigma_z^2}} + e^{\frac{-(z-2h+h_s)^2}{2\sigma_z^2}} + e^{\frac{-(z+2h-h_s)^2}{2\sigma_z^2}} + e^{\frac{-(z-2h-h_s)^2}{2\sigma_z^2}} \right) \quad (1)$$

The following assumptions are made regarding the Gaussian model: 1) The source is emitting CH₄ at a constant rate; 2) The mass of CH₄ is conserved when reflected at the surface of the ocean or the top of the boundary layer; 3) Wind speed and vertical eddy diffusivity are constant with time; 4) There is uniform vertical mixing; and 5) Terrain (ocean surface) is relatively flat between source and detector. The PGSC were determined for an offshore flow of air following the parametrizations described in Erbrink and Scholten (1995), Hanna et al. (1985) and Hsu (1992).”

At P6 L1:

“The height of the boundary layer is calculated from the Global Forecast System’s global forecast model archives (GFS, 2019).”

At P6 L10

“2.5 Uncertainties

Of the Gaussian plume model assumptions presented in Section 2.3, two may not be valid - uniform vertical mixing and a constant wind speed. The uncertainty in uniform vertical mixing is discussed in Section 3.4. To investigate how uncertainties in the measurements and modelling affect the calculated emission, we ran Gaussian plume model scenarios using data that reflect the input values’ uncertainty bounds. The scenarios run using the Gaussian plume approach were: varying wind speed (based on measurement); UGGA precision (± 2 ppb); thermometer precision (± 0.1 °C); the PGSC ($+ 1$ PGSC); and distance from detector to emission source (± 50 m). The uncertainties of the UGGA and thermometer were taken from literature. The uncertainty in the PGSC used reflects the possibility that the temperature of the natural gas leaving the subsurface could be hotter than air and therefore less stable. The uncertainty in distance from the emission source to the detector results from not knowing where gas is leaking; here we assume the leak could be from anywhere on a production platform that is 100 m long.”

P9 L10

“3.4 Uncertainties/shortcomings of Gaussian Plume modelling

A range of scenarios were run using the Gaussian plume model to estimate uncertainty in average CH₄ emissions resulting from UGGA instrument precision, thermometer precision, varying wind speed, assessment of the PGSC and uncertainty in distance between the emission source and the detector. Uncertainty in the UGGA and the thermometer have little effect on the average emission estimate (Supplementary Material Section 3). The largest variability in wind speed was recorded during measurement of platform # 3 on the 6th July 2017 at 4.4 ± 0.6 m s⁻¹ (Supplementary Material Section 1), using this variability in wind speed in the Gaussian plume model results in an uncertainty in average emission of ± 12 %. Uncertainty in estimating the distance between the emission source and the detector results in an uncertainty in average emissions of ± 8 %. The Gaussian plume model has the greatest response to the uncertainty in estimating the PGSC, resulting in an uncertainty of ± 41 %. We estimate the overall uncertainty in the average CH₄ emission, calculated as the root of the sum of the individual uncertainties squared, to be ± 45 %.

As mentioned in Section 2.5, the uniform vertical mixing assumption made in the Gaussian Plume model may not hold here as the data we collected provides no information on vertical mixing. However, the Gaussian plume model only assumes a constant vertical mixing rate between the source and the detector. In most cases this distance is relatively short and unlikely to significantly affect the calculation of emissions. In future experiments,

the vertical mixing rate could be calculated by measuring the vertical gradient of wind speeds to make an accurate thermodynamic profile.”

Critique #2

While the investigation that forms the bulk of the study is logically solid, the abstract begins with a line of reasoning that is quite circumstantial. It describes what motivated the study. While it is interesting to read about the authors motivation, excluding this information would make the definitive methods of the investigation clearer.

The passage is: “Recent studies suggest oil and natural gas production facilities in North America may be underestimating methane (CH₄) emissions during extraction. This, coupled with unusually high CH₄ mole fractions observed at coastal sites during onshore winds in the UK, suggests CH₄ emissions from oil and gas extractions in the North Sea could be higher than previously reported.”

I don’t think the conclusion necessarily follows. Emissions can vary greatly between facilities and across production fields. The geology and technology used in the North American fields where the aforementioned studies were conducted is much different than those of the North Sea. Unusually high mole fractions observed at the coast when winds came from the sea do not necessarily point to emissions from the oil and gas industry. Airmass trajectories can be quite complicated and there are many sources on a continental scale.

If the authors want to include the in-situ observations at Weymouth in the paper, then they should include a trajectory analysis. The paper does work without this passage, though.

Author’s response:

As suggested by the reviewer, using coastal observations to infer large emissions from offshore oil and gas operations is very difficult. The initial intention of the isotopic analysis of air collected during a Northerly wind event was to test if the air could have come from production platforms, i.e. has the isotopic signature as gas collected offshore. Unfortunately, as the analysis progressed it became apparent that these measurements could not be used to definitively identify the source.

To address these concerns in the manuscript we have removed the onshore/isotopic observations, the first sentence in the abstract has been removed and the second sentence edited.

Change to manuscript:

At P1 L17:

“**To investigate whether offshore oil and gas platforms leak methane (CH₄) during normal operation,** we measured CH₄ mole fractions around eight oil and gas production platforms in the North Sea which were neither flaring gas nor off-loading oil.”

Critique #3

What is the uncertainty of the parameters of the linear model from your Keeling plot (Figure 2a)? What is the uncertainty in the source isotopic signature? From the appearance of the plot, there is a very poor correlation between observations and very high uncertainty in the source isotopic signature. It might be instructive to color the points by time. The data are likely insufficient to describe the source. It could just be that all the data were taken near each other in time, and so there is not enough variation in the CH₄ concentration to extract a signal.

Author's response:

The reviewer's concerns over the isotopic analysis are justified. We feel that while this analysis could indicate offshore leakage, it does not necessarily compliment the central message of the manuscript, methane leaks from all offshore platforms. Instead of attempting to further justify the inclusion of the isotopic analysis to this body of work, we have decided to remove the section completely.

Critique #4

The description of the uncertainty analysis for the Gaussian plume model is lacking in detail, and there are some red flags. For one, the total uncertainty is given as +/-54% while the uncertainty due to stability class uncertainty alone is estimated at 54%, and the greatest source of uncertainty is said to be the emission height.

The uncertainty analysis does not explicitly define what is meant by "uncertainty". It is said "The overall uncertainty, calculated as the root of the sum of individual uncertainties squared...". This implies that the uncertainties are standard deviations of normally distributed random errors. But the uncertainties are almost certainly correlated.

Author's response:

New sections have been added to present and discuss the uncertainties in Gaussian Plume modelling.

Changes to the manuscript:

P6 L10

"2.5 Uncertainties

Of the Gaussian plume model assumptions presented in Section 2.3, two may not be valid - uniform vertical mixing and a constant wind speed. The uncertainty in uniform vertical mixing is discussed in Section 3.4. To investigate how uncertainties in the measurements and modelling affect the calculated emission, we ran Gaussian plume model scenarios using data that reflect the input values' uncertainty bounds. The scenarios run using the Gaussian plume approach were: varying wind speed (based on measurement); UGGA precision (± 2 ppb); thermometer precision (± 0.1 °C); the PGSC (+ 1 PGSC); and distance from detector to emission source (± 50 m). The uncertainties of the UGGA and thermometer were taken from literature. The uncertainty in the PGSC used reflects the possibility that the temperature of the natural gas leaving the subsurface could be hotter than air and therefore less stable. The uncertainty in distance from the emission source to the detector results from not knowing where gas is leaking; here we assume the leak could be from anywhere on a production platform that is 100 m long."

Technical correction #1

Page 1 The abstract should include a concise description of the methods

Author's reply:

A concise description of the methods has been included in the abstract

Change to manuscript:

P1 L19:

“We use the measurements from summer 2017, along with meteorological data, in a Gaussian plume model to estimate CH₄ emissions from each platform.”

Technical correction #2

Page 2, Figure 1 Caption: Mention Weymouth observatory.

Author’s reply:

The caption in Figure 1 has been edited

Change to manuscript:

P2 L14:

“The map also shows the location of the University of East Anglia’s Weybourne Atmospheric Observatory (WAO; 52.95 °N, 1.14 °E) in Weybourne, Norfolk.”

Technical correction #3

Page 3, Line 16: “To investigate the loss of CH₄ from offshore oil and gas installations we use two approaches; 1) determine whether the source of CH₄ enhancements at WAO could be from oil and gas production platforms; and 2) estimate an average CH₄ loss from offshore installations by making direct measurements of CH₄ emissions from off-shore production platforms in the North Sea”. The listed items don’t seem to be “approaches” to investigating the loss of CH₄. Is this a typographical error? It would be nice to see this information replaced with a clear and concise description of the methods used in the paper.

Author’s reply:

As suggested a brief description of the methods has been included.

Changes to manuscript:

P3 L23:

“To investigate the CH₄ loss from offshore oil and gas installations in UK waters, we measure CH₄ mixing ratios downwind of offshore platforms and use these data in a Gaussian plume model to estimate CH₄ emission rates. The CH₄ loss is then presented as a percentage of the CH₄ produced by each platform.”

Technical correction #4

Page 3, Line 24: “between 10:00 and 13:00” please include time zone.

Author’s response:

This section has been removed

Technical correction #5

Page 4, Line 6: “Measurements from boats of CH₄ emissions from individual production platforms...” Careful, the CH₄ mole ratio was measured and the emission rate was estimated using a simple model. I think it’s a reach to say that the emissions were measured. The previous sentence uses my preferred language.

Author’s reply:

Corrected as suggested.

Change to manuscript:

P3 L30

“Measurements were made during normal operation (i.e. pilot light on the flare stack was lit, but no flaring or offshore oil loading was observed)”

Technical correction #6

Page 5, Line 7: “The gas is considered to be well-mixed within the volume of the cone” This is an inaccurate description of the Gaussian Plume model. A Gaussian Plume describes the distribution of the mass of the gas at a given time as a multivariate Gaussian in space. To say that the gas is well mixed within a volume would suggest a uniform distribution in a finite region.

Author’s reply:

As suggested this line has been removed.

Technical correction #6

Page 6, Figure 2a): The correlation here looks very weak.

Author’s reply:

As the reviewer notes, the results from this measurement are ambiguous. To address this we have removed the section.

Technical correction #7

Page 8, Figure 3: Can this figure this be a plate showing all the observations rather than just the observations from 1 platform? I’m assuming the arrow shows the average wind speed and direction? How much variability was there?

Author’s reply:

All observations have been added as Supplementary Material Section 2.

During the measurements the highest uncertainty in wind speed during measurement was measured at $\pm 0.6 \text{ m s}^{-1}$ during the measurement of platform ID # 3 on the 6th July 2017.

P9 L14

“The largest variability in wind speed was recorded during measurement of platform # 3 on the 6th July 2017 at $4.4 \pm 0.6 \text{ m s}^{-1}$ (Supplementary Material Section 1), using this variability in wind speed in the Gaussian plume model results in an uncertainty in average emission of $\pm 12 \%$.”

Technical correction #8

Page 9, Line 4: “The main uncertainty using the Gaussian plume approach in this study is in estimating the height of emission...” I would argue that there are many large sources of uncertainty in the Gaussian plume model, some of which are almost certainly greater than error in the emission height. For example, the assumption of homogenous diffusion.

Author’s reply:

As suggested by the reviewer the section describing the uncertainties has been changed to include a systematic estimate of major uncertainties.

Changes to the manuscript:

P9 L10

“3.4 Uncertainties/shortcomings of Gaussian Plume modelling

A range of scenarios were run using the Gaussian plume model to estimate uncertainty in average CH₄ emissions resulting from UGGA instrument precision, thermometer precision, varying wind speed, assessment of the PGSC and uncertainty in distance between the emission source and the detector. Uncertainty in the UGGA and the thermometer have little effect on the average emission estimate (Supplementary Material Section 3). The largest variability in wind speed was recorded during measurement of platform # 3 on the 6th July 2017 at $4.4 \pm 0.6 \text{ m s}^{-1}$ (Supplementary Material Section 1), using this variability in wind speed in the Gaussian plume model results in an uncertainty in average emission of $\pm 12 \%$. Uncertainty in estimating the distance between the emission source and the detector results in an uncertainty in average emissions of $\pm 8 \%$. The Gaussian plume model has the greatest response to the uncertainty in estimating the PGSC, resulting in an uncertainty of $\pm 41 \%$. We estimate the overall uncertainty in the average CH₄ emission, calculated as the root of the sum of the individual uncertainties squared, to be $\pm 45 \%$.

As mentioned in Section 2.5, the uniform vertical mixing assumption made in the Gaussian Plume model may not hold here as the data we collected provides no information on vertical mixing. However, the Gaussian plume model only assumes a constant vertical mixing rate between the source and the detector. In most cases this distance is relatively short and unlikely to significantly affect the calculation of emissions. In future experiments, the vertical mixing rate could be calculated by measuring the vertical gradient of wind speeds to make an accurate thermodynamic profile.”

P10 L17

“The emission estimates presented here are from a pilot study and further work is needed to establish total CH₄ leakage rates from offshore oil and gas platforms. We have established, however, that CH₄ enhancements can be detected downwind of all production platforms during normal operations when neither venting, flaring or oil loading activities are taking place. Our measurements used in a Gaussian plume model indicate leakage from offshore installations are likely larger than previously estimated. However, these emission estimates come with large uncertainties as they are based on relatively few measured platforms, assume values for the height of emission, lateral and vertical mixing ratio distributions, and may not meet all the Gaussian plume model assumptions.

When the CH₄ emissions are calculated for two different emission heights, the importance of identifying the source location and height above the sea becomes apparent. The median CH₄ emissions from the five platforms is 6.8 g s^{-1} when the emissions are all assumed to come from the working deck, while the median emissions is $2,658 \text{ g s}^{-1}$ (47% of production) when all CH₄ is assumed to be emitted from the flare i.e. the highest point of the platform. This analysis indicates that the median emission presented here, based on the assumption that the

emissions occur from the working deck, is a conservative estimate. However, without further measurements the height of the emission source cannot be definitively determined and this leaves the possibility that leakage is higher during normal operations than our results indicate. The other input variables that cannot be determined without further measurement are the lateral and vertical mixing ratio distributions but we feel that following the study of Blackall et al. (2007) the estimates used in this study are sufficient to establish leakage from oil and gas platforms and to provide a rough estimate of their emissions. As with the emission height, mixing can be resolved with further measurement, including the use of aircraft to resolve the vertical and horizontal mixing of the plume. It is clear that further studies are needed to provide additional data that will yield more definitive emission estimates. Using the near-source (< 1 km) observations of this paper (Fig. 2; Supplementary Material Section 2 platform #5 and #6) we can see that plumes from the leaks are compact (<200 m wide) and in some cases difficult to detect from sea-level measurements (Supplementary Material Section 2 platforms #7 and #8). Making 3-dimensional observations downwind of the platforms and using a sonic anemometer would help identify some of the unknowns presented here. Also, measuring more platforms over a longer time frame would improve the understanding of ambient leakage.

Any further measurements would be significantly easier with the cooperation of the oil and gas industry which could benefit from the findings. If the emissions are as low as the industry currently estimates, further measurements confirming low leakage rates would improve consumer confidence in oil and gas extraction activities. Alternately, if emissions are higher than currently reported, additional measurements would give the industry an opportunity to identify common issues such as incomplete combustion at the flare (Fig. 2), reduce leakage, and improve the efficiency of platforms thus potentially increasing profits from the extracted gas.”

Reviewer: Daniel Varon

Specific comment #1

Page 2, Lines 13-14: To my knowledge, the studies cited (Zavala-Araiza et al., 2015 and Schwietzke et al., 2017) do not directly discuss methane emissions from offshore platforms. Perhaps this sentence could be broken into two sentences or parts, the first citing these studies as evidence that public inventories often underestimate methane emissions, and the second suggesting that the same may be true for offshore oil and gas platforms.

Author's response:

As suggested this sentence has been edited.

Change to manuscript:

P2 L19

“However, recent studies indicate public inventories in the United States underestimate CH₄ emissions including from the oil and gas supply chain (Alvarez et al., 2018; Zavala-Araiza et al., 2015, Schwietzke et al., 2017). This leads to the question: Could CH₄ emissions from offshore oil and gas platforms be higher than previously estimated?”

Specific comment #2

P2, L17-19: Nara et al. (2014) quantified methane emissions from offshore platforms in Southeast Asia using a mass balance approach, but the authors describe that study as qualitative rather than quantitative. It would be helpful to clarify this comparison in the manuscript.

Author's reply:

As suggested a comparison between the findings of this study and Nara et al. (2014) have been made and included in the introduction and the discussion.

Changes to manuscript:

P3 L6

“However, a mass-balance approach identifies CH₄ emissions from offshore oil and gas operations off the coast of South East Asia as having a large regional median (range) emission of 99 (4 – 427) g CH₄ s⁻¹ platform⁻¹ for the Malay Peninsula and 15 (2 – 46) g CH₄ s⁻¹ platform⁻¹ for Borneo (Nara et al., 2015).”

P11 L29

“Moreover, the median value of this study (6.8 g s⁻¹) is much smaller than the regional median emission estimate of 99 g s⁻¹ for the Malay Peninsula and 15 g s⁻¹ for Borneo (Nara et al., 2015), which suggests that the ambient leakage rate may be lower in the North Sea than other regions of the world.”

Specific comment #3

P3, L19-20: I would recommend removing this novelty claim, because the study is clearly original. Targeted measurement of methane emissions from individual oil and gas platforms is an impressive contribution. This sentence could be replaced with a one- or two-sentence comparison to the previous work of Nara et al. (2014).

Author's comment:

Thank you for the endorsement of our work. As suggested, we have removed the sentence about novelty.

Specific comment #4

P4, L24: The maximum horizontal distance from the platforms is reported to be 1500 m, but some platforms in Table 1 have distances of 2000 m.

Author's reply:

This was a typo and has been corrected.

Change to manuscript:

P4 L21

“and 2 km horizontal distance”

Specific comment #5

P8, Figure 3: Peak enhancements (2160-2230 ppb) do not match the value reported in Table 1 (2290 ppb). Can the authors clarify in the table caption (or elsewhere in the manuscript) whether the downwind methane concentrations reported in Table 1 represent peak concentrations, or something else?

Author's response:

This should have been platform ID #6 instead of # 5. The caption has been corrected.

Change to manuscript:

P8 Caption Figure 2

“Minute-averaged CH₄ mole fraction measurements made upwind and downwind of production platform, ID # 6, on the 24th of August.”

Specific comment #6

P8, L11-14: The total emission from the 8 platforms should not be compared to the total production from only 6 platforms unless there is good reason to believe that the missing production rates are small. Indeed, if one of platforms #1 or #2 produced as much gas as platform #4, the calculation would be quite different. One solution to this problem would be to compare emissions and production rates only for platforms #3-#8. Another option would be to impute the production rates for platforms #1 and #2 from the average (or median) of the other platforms' rates.

Author's response:

As suggested we have changed the calculation to only include platforms 3 to 8. Text has been included to the caption of Table 1 and the manuscript to reflect this.

Change to manuscript:

P7 Table 1 caption

“The calculation of the “Median”, “Mean” and “Total” only use data from platforms #4 to # 8. Platforms #1 and #2 did not have production data available for the time of measurement. During the measurement of Platform #3 the height of the PBL was calculated as zero (GFS, 2019) making the Gaussian plume modelled emission estimate ambiguous.”

P8 L11

“During the measurement of platform #3, the calculated boundary layer height was 0 m (GFS, 2019) making the emission estimate ambiguous and, even though presented in Table 1, has not been used further in the analysis. Using emission data from the five platforms with available production data and with a non-zero calculated PBL (platforms #4 through #8), the median CH₄ emission was 6.8 g s⁻¹ (mean 11.2 g s⁻¹).”

Specific comment #7

P9, L12-15: Why might the Pasquill-Gifford stability classes used to infer emissions from the platforms be too stable? What would cause the difference between stability at the receptor and stability at the source? Is it the difference in wind speed between the surface and 40-90 m altitude? If so, would this not suggest that the stability class as assessed at the surface might be too unstable (due to the winds being faster at altitude)? One additional sentence would probably clear this up.

Author’s response:

The methane lost from the platform may be less stable as it has come from the subsurface and may be a warmer than the surrounding air and therefore less stable. As a test, we suggest this could be 1 PGSC less stable than calculated. To clarify this text has been added.

Change to manuscript:

P6 L11

“The uncertainty in the PGSC used reflects the possibility that the temperature of the natural gas leaving the subsurface could be hotter than air and therefore less stable.”

Specific comment #8

P10, L8: Why are the estimated platform emissions larger than BEIS reported emissions of 0.13% by a factor of 2, but similar in magnitude to NAEI emissions? From page 6, line 1, it seems like the BEIS and NAEI figures should be similar, since the BEIS data “form the basis for emissions reported under category 1B2 within the National Atmospheric Emissions Inventory (NAEI; BEIS, 2018).” This can also probably be clarified in a sentence.

Author’s reply:

The platform emissions are twice as large as the BEIS emission estimates but appear to be consistent with the NAEI because NAEI currently only accounts for venting and flaring not leakage. Here we present the leakage estimates only as venting and flaring were not taking place. It is only by coincidence that our leakage estimates are the same as the NAEI values.

Change to manuscript:

P10 L11

“neither of which was taking place during our measurements”

Specific comment #9

P10, L25-31: I am a bit hesitant to draw broad conclusions about global methane emissions from the oil and gas sector based on results from a small number of offshore platforms. It is interesting that the Oil and Gas Climate Initiative does not include ambient emissions in its global estimates when these emissions seem to be significant (as the authors illustrate), but I would expect their magnitude to vary greatly across geographies and industries. Indeed, the authors make note of this variability on page 2, line 15, and mention also the particularly harsh environment of the North Sea on page 10, line 20. I would recommend that the authors more

clearly qualify their extrapolation of ambient emissions from North Sea offshore platforms to ambient emissions from global oil and gas activities.

Author's reply:

The text has been amended to reflect the speculative nature of this statement. The idea of this paragraph was to merely represent the concept of emissions from leakage and the potential impact of these measurements.

Change to manuscript:

P11 L25

“If a global CH₄ emission from ambient leakage of 0.19% estimated by this study (0.8 Tg CH₄ yr⁻¹) is added to the current global estimate from flaring, venting and offshore oil loading (1.6 Tg CH₄ yr⁻¹) the total CH₄ emission from offshore oil and gas production would increase significantly. It should be noted that the value of 0.19% is based on a very small sample size using a method that comes with significant uncertainty.”

Technical correction #1

Page 1, Line 4: The words “onshore” and “offshore” are spelled differently throughout the text, both with and without dashes.

Author's reply:

Have amended to be consistently “onshore” and “offshore”.

Technical correction #2

P2, L10: The acronym “OGA” is not defined.

Author's reply

The acronym has been defined as the UK Oil and Gas Authority.

Change to MS:

P2 L16:

“(UK Oil and Gas Authority, 2018).”

Technical correction #3

P3, L11: The acronym “EEMS” is not defined.

Author's reply:

EEMS has been defined as the Environmental and Emissions Monitoring System.

Change to MS:

P3 L17:

“UK Government's Department of Energy and Climate Change Environmental and Emissions Monitoring System (DECC EEMS, 2008)”

Technical correction #4

P7, L10-11: Redundant use of the word “example.”

Author's reply:

Deleted as suggested.

Technical correction #5

P9, L8: It seems like there might be a missing word here.

Author's reply:

Have amended the sentence.

Change to MS:

P8 L14

“As a sensitivity study, the median modelled emission is $2,658 \text{ g s}^{-1}$ (mean $1,892 \text{ g s}^{-1}$) when we assume all CH_4 is emitted from the highest point of the platform, i.e. the flare.”

Methane emissions from oil and gas platforms in the North Sea

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Abstract.

Since 1850 the concentration of atmospheric methane (CH₄), a potent greenhouse gas, has more than doubled. Recent studies suggest that emission inventories may be missing sources and underestimating emissions. To investigate whether offshore oil and gas platforms leak CH₄ during normal operation, we measured CH₄ mole fractions around eight oil and gas production platforms in the North Sea which were neither flaring gas nor off-loading oil. We use the measurements from summer 2017, along with meteorological data, in a Gaussian plume model to estimate CH₄ emissions from each platform. We find CH₄ mole fractions of between 11 and 370 ppb above background concentrations downwind of the platforms measured, corresponding to a median CH₄ emission of 6.8 g CH₄ s⁻¹ for each platform, with a range of 2.9 to 22.3 g CH₄ s⁻¹. When matched to production records, during our measurements individual platforms lost between 0.04% and 1.4% of gas produced with a median loss of 0.23%. When the measured platforms are considered collectively, (i.e. the sum of platforms' emission fluxes weighted by the sum of the platforms' production), we estimate the CH₄ loss to be 0.19% of gas production. These estimates are substantially higher than the emissions most recently reported to the National Atmospheric Emission Inventory (NAEI) for total CH₄ loss from United Kingdom platforms in the North Sea. The NAEI reports CH₄ losses from the offshore oil and gas platforms we measured to be 0.13% of gas production, with most of their emissions coming from gas flaring and offshore oil loading, neither of which were taking place at the time of our measurements. All oil and gas platforms we observed were found to leak CH₄ during normal operation and much of this leakage has not been included in UK emission inventories. Further research is required to accurately determine total CH₄ leakage from all offshore oil and gas operations and to properly include the leakage in national and international emission inventories.

1 Introduction

Methane (CH₄) is a greenhouse as well as a precursor of tropospheric ozone, which is widely regulated as a component of photochemical smog. Since 1850 atmospheric CH₄ mixing ratios have increased from 715 ppb to 1865 ppb in February 2019 with an annual increase of 10ppb/year in 2018 (NOAA, 2019). This increase is largely driven by anthropogenic activities though uncertainties exist in the magnitude of individual source sectors and

sinks (Turner et al., 2019). Observatories have been set up around the world to track trends in CH₄ concentrations (de Coninck et al., 2018).

Between 2012 and 2015 unusually high CH₄ enhancements of up to 400 ppb above background were observed at the University of East Anglia's Weybourne Atmospheric Observatory (WAO; 52.95 °N, 1.14 °E) during periods of northerly onshore winds and high surface pressures (Connors, 2015; Staunton-Sykes, 2016; Connors et al., 2018). These elevated enhancements were unexpected as the air came from the open ocean. However, a potential source of these CH₄ enhancements is leakage from offshore oil and gas production platforms 80 km away from WAO in the North Sea (OSPAR, 2018; Fig. 1).

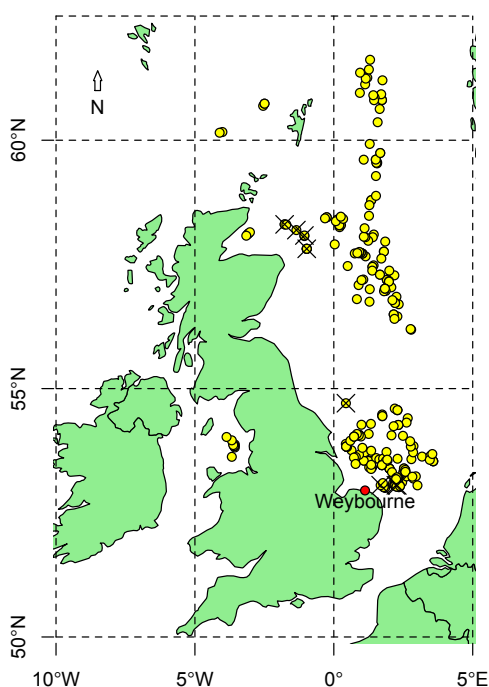


Figure 1 Map of the North Sea showing the locations of all UK offshore oil/gas platforms (the filled yellow circles) and the eight platforms measured by this study (black crosses) (Source: OSPAR, 2012). **The map also shows the location of the University of East Anglia's Weybourne Atmospheric Observatory (WAO; 52.95 °N, 1.14 °E) in Weybourne, Norfolk, UK.**

In 2015 the UK extracted about 32 Tg of natural gas from the North Sea (UK Oil and Gas Authority, 2018). During this time, a loss of 40 Gg CH₄ (0.13 % of natural gas production) was reported by the UK Government Department for Business, Energy & Industrial Strategy (BEIS), mainly through venting (24 Gg CH₄ yr⁻¹) and flaring (12 Gg CH₄ yr⁻¹) activities (BEIS, 2018). However, recent studies indicate public inventories in the United States underestimate CH₄ emissions including from the oil and gas supply chain (Alvarez et al., 2018; Zavala-Araiza et al., 2015, Schwietzke et al., 2017). **This leads to the question: Could CH₄ emissions from offshore oil and gas platforms be higher than previously estimated?**

Land-based measurements in West Virginia and Colorado, USA, estimate that onshore oil and gas extraction activities lose between 0.1 and 10 % of CH₄ produced (Petron et al., 2012, Omara et al., 2016; Schwietzke et al., 2017; Alvarez et al., 2018; Englander et al., 2018; Riddick et al., 2019). At present there are no direct, near-source measurements of CH₄ emissions from offshore oil and gas production. **However, a mass-balance approach identifies CH₄ emissions from offshore oil and gas operations off the coast of South East Asia as having a large regional median (range) emission of 99 (4 – 427) g CH₄ s⁻¹ platform⁻¹ for the Malay Peninsula and 15 (2 – 46) g**

$\text{CH}_4 \text{ s}^{-1} \text{ platform}^{-1}$ for Borneo (Nara et al., 2015). Numerous productive offshore oil and gas fields exist across the globe, including Saudi Arabia, Brazil, Mexico, Norway and the United States, making careful measurement and analysis of leakage from these platforms important for global emission inventories.

Several activities on offshore production platforms in the North Sea are explicitly identified by the BEIS in the National Atmospheric Emission Inventory (NAEI) as sources of CH_4 emission including: combustion activities such as gas flaring, offshore oil loading and venting directly to the atmosphere (BEIS, 2018). Leakage during normal operations are not explicitly included. Oil and gas operators report an annual CH_4 emission estimate for each offshore production platform to the NAEI; these emission estimates are primarily calculated using emission factors (Butterfield, 2017). Technical guidance on the emission factor based calculations is available through the UK Government's Department of Energy and Climate Change **Environmental and Emissions Monitoring System** (DECC EEMS, 2008). The main shortcoming of using emission factors and activity levels to estimate total emissions is that total emissions can be underestimated if not all emission sources are identified. For example, leaks not obvious to platform personnel would not be included and the total emission would be an underestimate of CH_4 lost. Overall, as emission factor calculations rely on explicit knowledge of all sources of leakage, current approaches used by industry could underestimate total CH_4 emissions from offshore installations.

In this study we investigate CH_4 emissions from offshore oil and gas installations in the North Sea and determine how they differ from those currently reported by the BEIS. **To investigate the CH_4 loss from offshore oil and gas installations in UK waters, we measure CH_4 mixing ratios downwind of offshore platforms and use these data in a Gaussian plume model to estimate CH_4 emission rates. The CH_4 loss is then presented as a percentage of the CH_4 produced by each platform.**

2 Methods - Boat based observations

Oil and gas platforms in UK waters are located between 30 and 500 km from the UK mainland, with the majority of platforms located to the east of the UK in the North Sea (Fig. 1; OSPAR, 2018). To investigate possible emissions from these platforms, sea-level CH_4 mole fractions were measured around eight oil and gas platforms between the 6th June and 25th August 2017. **Measurements were made during normal operation** (i.e. pilot light on the flare stack was lit, but no flaring or offshore oil loading was observed). Where possible a full circle was made around the installation to observe the upwind and downwind methane mole fractions. To determine if flaring or offshore oil loading was occurring, a visual inspection was made of the installation. We assumed that venting was not taking place because no venting was reported in any of the most recent NAEI. Previously published emissions from the measured platforms in the NAEI are reported to be almost entirely due to flaring (83%) and offshore oil loading (17%), with reported emissions generated using emissions factors (Brown et al., 2017; BEIS, 2018).

The oil and gas platforms measured here were selected at random, constrained only by the need for accessibility. Fishing boats were chosen as the measurement platforms because of budgeting and availability constraints. Maritime and Coastguard Agency regulations for the available vessels (MCA category 2) meant that the platforms measured had to be less than 60 miles from a safe haven. Four of the eight platforms only produced natural gas (#1 to #4) that was transported to the mainland via pipeline, while the remaining four produced oil and gas. Two of the oil and gas platforms (#5 and #6) include floating production storage and offloading vessels, which receive hydrocarbons, process them and store them until they can be offloaded by tanker or pipeline, and the other two

platforms (#7 and #8) transport oil and gas directly to the mainland by pipeline. Methane mole fractions, latitude, longitude, and meteorological data were collected as the boat travelled upwind and downwind of the platforms.

2.1 Methane mole fraction measurements – Los Gatos UGGA

The Los Gatos Research Ultra-portable Greenhouse Gas Analyzer (UGGA; www.lgrinc.com) was used to measure CH₄ concentrations near the off-shore oil and gas platforms. The UGGA is a laser absorption spectrometer that measures CH₄ mole fractions in air (Paul et al., 2001). The UGGA reports CH₄ mole fractions every second, with a stated precision of < 2 ppb (1 σ @ 1 Hz) over an operating range of 0.1 to 100 ppm. Calibration of the UGGA was conducted before and after deployment using low (1.93 ppm CH₄), target (2.03 ppm CH₄) and high (2.74 ppm CH₄) mole fraction gases calibrated on the World Meteorological Organization (WMO) scale. Measurements were taken between the edge of the exclusion zone (500 m from the platform; HSE, 2018) and 2 km horizontal distance from the platforms. The inlet line was attached to a mast 2.5 m above sea level, to avoid contamination from the boat's exhaust, and protected from water incursion using an aluminium funnel. The air was filtered using a 2 μ m filter. Background CH₄ mole fractions were measured while the boat was upwind of the production platform.

2.2 Meteorological data

Meteorological data were collected using a wireless weather station (Maplin, UK) attached to a mast 2 m above sea level. Data were sampled and recorded at one-minute intervals and included: wind speed (u , m s⁻¹), wind direction (WD , ° to North), air temperature at 2 m (T_a , K), relative humidity (RH , %), rain rate (R , mm hr⁻¹), irradiance (I , W m⁻²) and air pressure (P , Pa). The wind speed used in the emission modelling was corrected for emission height using a wind profile power law (Touma, 1977; Hsu et al., 1994).

2.3 Gaussian Plume Model

The Gaussian plume model used in this study calculates the mole fraction of a gas as a function of distance downwind from a point source (Seinfeld and Pandis, 2006). As a gas is emitted, it is entrained in the prevailing ambient air flow and disperses in the y and z directions (relative to a mean horizontal flow in the x direction) with time, forming a cone. The mole fraction of the gas as a function of distance downwind depends on the emission flux at the source, the advective wind speed (u , m s⁻¹) and the rate of dispersion. The mole fraction of the gas (X , μ g m⁻³), at any point x meters downwind of the source, y meters laterally from the centre line of the plume, and z metres above ground level can be calculated (Eq. 1) using the source strength (Q , g s⁻¹), the height of the source (h_s , m), the height of the boundary layer (h , m) and the stability of the air (CERC, 2017; Hunt, 1982; Hunt et al., 1988). The standard deviation of the lateral (σ_y , m) and vertical (σ_z , m) mixing ratio distributions are calculated from the Pasquill Gifford stability class (PGSC) of the air (Pasquill, 1962; Busse and Zimmerman, 1973; US EPA, 1995). Even though this modelling method is relatively simple, offshore emissions estimates using the same parameterization of σ_y and σ_z were made by Blackall et al. (2008) and were in good agreement ($R^2 = 0.85$) with emissions calculated from a concurrent tracer release experiment. Alternative offshore parameterisations for σ_y and σ_z exist and are used in the EPA recommended Offshore and Coastal Dispersion Model (Hanna et al, 1985). However, these algorithms require further data on the micrometeorology which are not available and were therefore not used as they introduce additional unquantifiable uncertainty.

$$X(x, y, z) = \frac{Q}{2\pi u \sigma_y \sigma_z} e^{-\frac{y^2}{2\sigma_y^2}} \left(e^{-\frac{(z-h_s)^2}{2\sigma_z^2}} + e^{-\frac{(z+h_s)^2}{2\sigma_z^2}} + e^{-\frac{(z-2h+h_s)^2}{2\sigma_z^2}} + e^{-\frac{(z+2h-h_s)^2}{2\sigma_z^2}} + e^{-\frac{(z-2h-h_s)^2}{2\sigma_z^2}} \right) \quad (1)$$

The following assumptions are made regarding the Gaussian model: 1) The source is emitting CH₄ at a constant rate; 2) The mass of CH₄ is conserved when reflected at the surface of the ocean or the top of the boundary layer; 3) Wind speed and vertical eddy diffusivity are constant with time; 4) There is uniform vertical mixing; and 5) Terrain (ocean surface) is relatively flat between source and detector. The PGSC were determined for an offshore flow of air following the parametrizations described in Erbrink and Scholten (1995), Hanna et al. (1985) and Hsu (1992).

2.4 Gaussian Plume model parameterization

A Gaussian plume approach was used to infer the CH₄ emissions flux from the oil and gas platforms using the CH₄ mole fraction data collected downwind. We used measurements for the mole fraction, and rearranging Equation 1 solved for the source term Q. Data used as input to the Gaussian plume model are: wind speed, wind direction, temperature, minute-averaged CH₄ mole fraction at 2 m, background CH₄ mole fraction and the PGSC. For the minute-averaged CH₄ mole fraction data, we assume the 1-minute averaged data near the centre of the observed instantaneous plume is representative of the centre of the time-averaged Gaussian plume. The PGSC are estimated from wind speed and irradiance data (Turner, 1970; Seinfeld and Pandis, 2006), as measured by the meteorological station on the boat. The height of the boundary layer is calculated from the Global Forecast System's global forecast model archives (GFS, 2019).

An unknown variable used in the Gaussian plume model in this study is the height at which emissions are released. The emissions could have come from the working deck of the platform, the top of the flare or somewhere in between. For the purposes of the emission estimates calculated and presented here, we assume CH₄ is emitted from the working deck only which results in the smallest emissions possible for a given measurement. As a sensitivity study, emissions were also calculated assuming the source was at the top of the flare stack only (see Supplementary Material Section 1). The height of the working deck and the height of the flare stack at each of the platforms were determined using platform characteristics data from each oil and gas platform available on the internet.

2.5 Uncertainties

Of the Gaussian plume model assumptions presented in Section 2.3, two may not be valid - uniform vertical mixing and a constant wind speed. The uncertainty in uniform vertical mixing is discussed in Section 3.4. To investigate how uncertainties in the measurements and modelling affect the calculated emission, we ran Gaussian plume model scenarios using data that reflect the input values' uncertainty bounds. The scenarios run using the Gaussian plume approach were: varying wind speed (based on measurement); UGGA precision (± 2 ppb); thermometer precision (± 0.1 °C); the PGSC (+ 1 PGSC); and distance from detector to emission source (± 50 m). The uncertainties of the UGGA and thermometer were taken from literature. The uncertainty in the PGSC used reflects the possibility that the temperature of the natural gas leaving the subsurface could be hotter than air and therefore less stable. The uncertainty in distance from the emission source to the detector results from not

knowing where gas is leaking; here we assume the leak could be from anywhere on a production platform that is 100 m long.

2.6 Data sources

The UK Department for Business, Energy & Industrial Strategy (BEIS) keeps the Environmental and Emissions Monitoring System (EEMS) which is the environmental database of the UK oil and gas industry. Methane emission data are uploaded to this by industry partners. These data form the basis for emissions reported under category 1B2 within the National Atmospheric Emissions Inventory (NAEI; BEIS, 2018). For details of how this data is incorporated into the NAEI, see Brown et al. (2017). The most recent point-source emission database from the NAEI available at the time of writing was for the year 2015. Individual platform production data for both 2015 and 2017 were taken from the Petroleum Production Reporting system published by the UK Oil and Gas Authority (OGA, 2018).

3 Results

3.1 Methane mole fractions around North Sea oil and gas platforms

Our sea-level surveys indicate CH₄ mole fraction enhancements can be measured near all of the production platforms observed, when upwind CH₄ mole fractions ($[CH_4]_{bgd}$, ppb) are compared with downwind mole fractions ($[CH_4]$, ppb; Table 1). The largest enhancement of 370 ppb was observed downwind of Platform #6 on the 24th August 2017, while the lowest enhancement of 11 ppb was observed downwind of platform #8 on the 25th August 2017. The median CH₄ enhancement downwind of the eight platforms was 43 ppb (mean: 112 ppb; range: 11 – 370 ppb). While measurements were being conducted, a maximum variability in wind speed of $\pm 0.6 \text{ m s}^{-1}$ was measured at platform #3 on the 6th July 2017; during no measurements did an observable change in wind direction occur. Complete circles of all installations were not possible due to access restrictions, i.e. the measurement vessel could not get between some platforms and maintain the 500 m clearance required of each platform, and there were occasions when the measurement boat was actively blocked by the platform's standby vessel (Supplementary Material Section 2).

Table 1. Meteorological, position and mole fraction data taken during the boat-based measurement campaign around oil and gas production platforms in the North Sea between 6th June and 25th August 2017. Emissions were calculated assuming the source of the emissions was at the working deck level. The calculation of the “Median”, “Mean” and “Total” only use data from platforms #4 to #8. Platforms #1 and #2 did not have production data available for the time of measurement. During the measurement of Platform #3 the height of the PBL was calculated as zero (GFS, 2019) making the Gaussian plume modelled emission estimate ambiguous.

ID #	Measurement Date	Type	Height (m)	Peak enhancement at the centre of the plume (ppb)	Emission (g s ⁻¹)	Platform CH ₄ production (g s ⁻¹)	Loss (%)
1	Jun 6	Gas	40	50	4.9	N/A	
2	Jun 6	Gas	40	51	4.9	N/A	
3	Jul 5	Gas	40	34	1.1	672	0.17
4	Aug 16	Gas	50	30	5.7	15,230	0.04
5	Aug 24	Oil/Gas	50	370	22.3	1,585	1.41
6	Aug 24	Oil/Gas	50	312	18.1	1,845	0.98
7	Aug 25	Oil/Gas	50	35	6.8	2,952	0.23
8	Aug 25	Oil/Gas	50	11	2.9	8,047	0.04
				Median (#4 - #8)	6.8		0.23
				Mean (#4 - #8)	11.2		0.54
				Total (#4 - #8)	55.8	29,662	
				Loss of CH ₄ produced (%) (#4 - #8)	0.19		

3.2 Source of leaks

Although the production platforms measured in this campaign were not actively flaring gas, (i.e. burning gas to reduce pressure during oil extraction), the pilot light on the top of the flare stack was actively burning gas. As an example, Fig.2 shows the minute-averaged CH₄ enhancements upwind and downwind of a production platform on the 24th August. This example was chosen as it was the only platform that had an offset flare stack (i.e. not centred in the platform). Fig.2 indicates the largest enhancement was downwind of the flare stack. The width of the plume (< 200 m) suggests a compact CH₄ source. This could be associated with incomplete combustion of natural gas feeding the pilot light at the top of the platform, or it could be associated with gas being emitted at the working deck level.

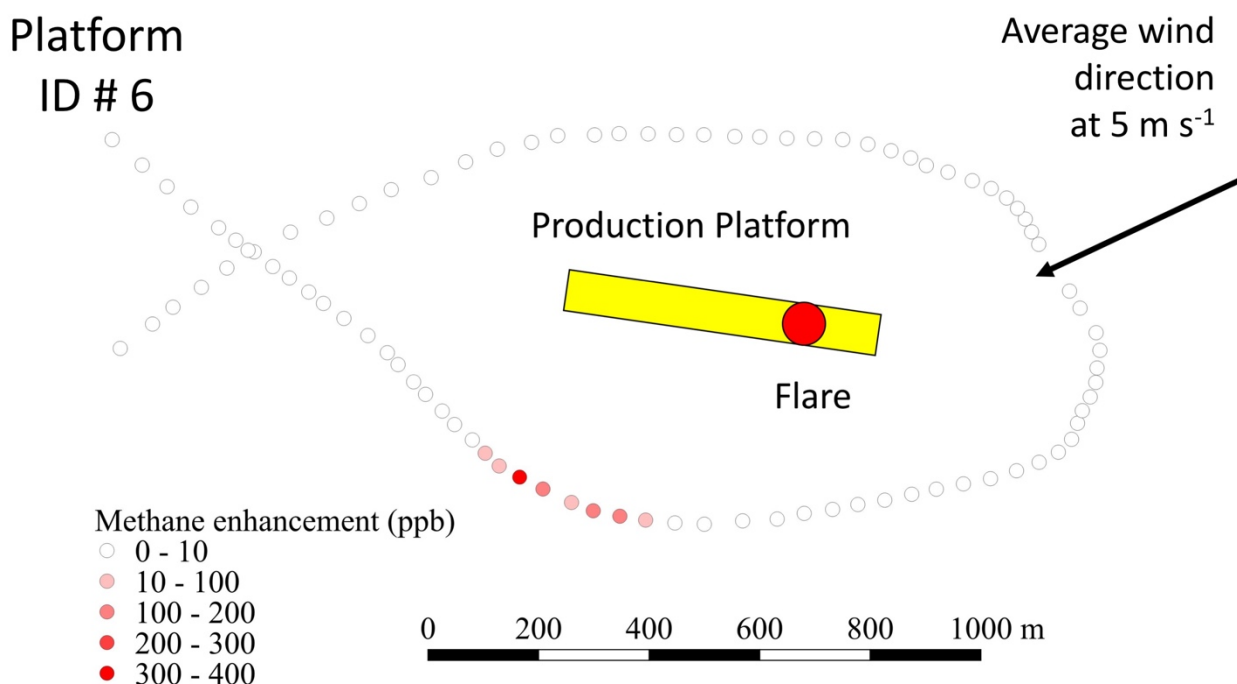


Figure 2. Minute-averaged CH₄ enhancements made upwind and downwind of production platform # 6, on the 24th of August 2017.

3.3 Estimating methane emissions

Using Gaussian plume modelling and assuming all emissions came from the working deck, the highest emission of 22.3 g s⁻¹ CH₄ was observed from platform # 5 on the 24th August 2017 while the lowest emission, 2.9 g s⁻¹, was observed on the 25th August 2017 from platform # 8. **During the measurement of platform #3, the calculated boundary layer height was 0 m (GFS, 2019) making the emission estimate ambiguous and, even though presented in Table 1, has not been used further in the analysis. Using emission data from the five platforms with available production data and with a non-zero calculated PBL (platforms #4 through #8), the median CH₄ emission was 6.8 g s⁻¹ (mean 11.2 g s⁻¹). As a sensitivity study, the median modelled emission is 2,658 g s⁻¹ (mean 1,892 g s⁻¹) when we assume all CH₄ is emitted from the highest point of the platform, i.e. the flare.**

When normalized against natural gas production data (OGA, 2018), the highest CH₄ loss rate corresponded to 1.4 % of production at platform #5 while the lowest loss rate corresponded to 0.04 % of production at platforms #4 and #8. We estimate the median CH₄ loss from platforms #4 through #8 to be 0.23 % of production. When weighted by production, i.e. the collective emission from the measured platforms (56 g s⁻¹; Table 1) as a fraction of the collective production of the measured platforms (29,662 g s⁻¹; Table 1), the average loss from all measured platforms was 0.19 % of their total production.

For comparison, we have also calculated the reported loss rates for 2015 using the most recent NAEI emissions data (Brown et al., 2017; BEIS, 2018). We find the median reported loss rate from NAEI was 0.23 % for the six platforms we measured where production data was available, with a production-weighted average of 0.19 %. These values are close to those we calculated. However, this apparent consistency is misleading as the NAEI emissions are dominated by CH₄ emissions from flaring and offshore oil loading activities neither of which were occurring during our measurement periods; this is discussed further in section 4.

3.4 Uncertainties/shortcomings of Gaussian Plume modelling

A range of scenarios were run using the Gaussian plume model to estimate uncertainty in average CH₄ emissions resulting from UGGA instrument precision, thermometer precision, varying wind speed, assessment of the PGSC and uncertainty in distance between the emission source and the detector. Uncertainty in the UGGA and the thermometer have little effect on the average emission estimate (Supplementary Material Section 3). The largest variability in wind speed was recorded during measurement of platform # 3 on the 6th July 2017 at $4.4 \pm 0.6 \text{ m s}^{-1}$ (Supplementary Material Section 1), using this variability in wind speed in the Gaussian plume model results in an uncertainty in average emission of $\pm 12 \%$. Uncertainty in estimating the distance between the emission source and the detector results in an uncertainty in average emissions of $\pm 8 \%$. The Gaussian plume model has the greatest response to the uncertainty in estimating the PGSC, resulting in an uncertainty of $\pm 41 \%$. We estimate the overall uncertainty in the average CH₄ emission, calculated as the root of the sum of the individual uncertainties squared, to be $\pm 45 \%$.

As mentioned in Section 2.5, the uniform vertical mixing assumption made in the Gaussian Plume model may not hold here as the data we collected provides no information on vertical mixing. However, the Gaussian plume model only assumes a constant vertical mixing rate between the source and the detector. In most cases this distance is relatively short and unlikely to significantly affect the calculation of emissions. In future experiments, the vertical mixing rate could be calculated by measuring the vertical gradient of wind speeds to make an accurate thermodynamic profile.

4 Discussion

From boat based observations we observed elevated CH₄ mole fractions, between 11 and 370 ppb above background, downwind of eight oil and gas production platforms in the North Sea when none of the platforms was engaged in either gas flaring or oil transfer and unloading. This suggests that all observed oil and gas platforms leak CH₄ during normal operations. Using the near-source CH₄ measurements in a simple Gaussian plume model (where the CH₄ emissions are calculated from the minute-averaged peak enhancement at the centre of the plume), we found the median of the calculated CH₄ emissions from offshore oil and gas installations to be $589 \text{ kg CH}_4 \text{ day}^{-1}$, with individual platform's CH₄ emissions ranging from 98 to $1,928 \text{ kg CH}_4 \text{ day}^{-1}$. Matching production data to our measurements we estimate 1) a median loss of CH₄ from the six platforms, unweighted by production, of 0.23 % (mean 0.54 %) and 2) the cumulative loss of CH₄, weighted by total production, of 0.19%. These results indicate that, of the platforms measured, those producing more gas leaked proportionally less of what they produced. Also, the two higher emitting platforms (#5 and #6) include floating production storage and offloading vessels; we find these to have much larger loss rates than the three fixed platforms (#4, #7 and #8). However, we also acknowledge our sample size is small and the five platforms may not be indicative of the overall performance of platforms in the North Sea.

The 2015 emission factor based NAEI emissions are within the ranges calculated in this study, i.e. a median loss rate of 0.23% and a production-weighted loss of 0.19%, and also show larger losses come from lower producing platforms. However, the NAEI provides the main source of emission for each installation and their reported emissions from the six platforms are almost entirely due to flaring (83%) and offshore oil loading (17%), **neither of which was taking place during our measurements**. Typically, these activities are not continuous on North Sea

platforms; consequently, emission rates are likely to be much higher at certain times than others. As flaring and oil loading did not coincide with our measurement campaign, the measured emissions presented here represent leakage only and do not account for intermittent emissions due to venting, flaring or oil loading activities. This suggests a potentially large missing source of CH₄ emissions in the national U.K. CH₄ emission inventory.

The emission estimates presented here are from a pilot study and further work is needed to establish total CH₄ leakage rates from offshore oil and gas platforms. We have established, however, that CH₄ enhancements can be detected downwind of all production platforms during normal operations when neither venting, flaring or oil loading activities are taking place. Our measurements used in a Gaussian plume model indicate leakage from offshore installations are likely larger than previously estimated. However, these emission estimates come with large uncertainties as they are based on relatively few measured platforms, assume values for the height of emission, lateral and vertical mixing ratio distributions, and may not meet all the Gaussian plume model assumptions.

When the CH₄ emissions are calculated for two different emission heights, the importance of identifying the source location and height above the sea becomes apparent. The median CH₄ emissions from the five platforms is 6.8 g s⁻¹ when the emissions are all assumed to come from the working deck, while the median emissions is 2,658 g s⁻¹ (47% of production) when all CH₄ is assumed to be emitted from the flare i.e. the highest point of the platform. This analysis indicates that the median emission presented here, based on the assumption that the emissions occur from the working deck, is a conservative estimate. However, without further measurements the height of the emission source cannot be definitively determined and this leaves the possibility that leakage is higher during normal operations than our results indicate. The other input variables that cannot be determined without further measurement are the lateral and vertical mixing ratio distributions but we feel that following the study of Blackall et al. (2007) the estimates used in this study are sufficient to establish leakage from oil and gas platforms and to provide a rough estimate of their emissions. As with the emission height, mixing can be resolved with further measurement, including the use of aircraft to resolve the vertical and horizontal mixing of the plume. It is clear that further studies are needed to provide additional data that will yield more definitive emission estimates. Using the near-source (< 1 km) observations of this paper (Fig. 2; Supplementary Material Section 2 platform #5 and #6) we can see that plumes from the leaks are compact (<200 m wide) and in some cases difficult to detect from sea-level measurements (Supplementary Material Section 2 platforms #7 and #8). Making 3-dimensional observations downwind of the platforms and using a sonic anemometer would help identify some of the unknowns presented here. Also, measuring more platforms over a longer time frame would improve the understanding of ambient leakage.

Any further measurements would be significantly easier with the cooperation of the oil and gas industry which could benefit from the findings. If the emissions are as low as the industry currently estimates, further measurements confirming low leakage rates would improve consumer confidence in oil and gas extraction activities. Alternately, if emissions are higher than currently reported, additional measurements would give the industry an opportunity to identify common issues such as incomplete combustion at the flare (Fig. 2), reduce leakage, and improve the efficiency of platforms thus potentially increasing profits from the extracted gas.

The continuous leakage of CH₄ from offshore production platforms observed here is consistent with observations of similar onshore operations (Omara et al., 2016; Riddick et al., 2019). Ambient leakage is not unexpected as these offshore production platforms are located in the inhospitable conditions of the North Sea, where wind speeds

regularly exceed hurricane force and waves can reach the working deck. However, it is surprising that ambient leakage has not been explicitly factored into the UK national emissions inventory, which relies solely on operators self-reported emissions calculated using emission factors combined with specific processes like flaring. Without direct measurement, operators can remain unaware of small emissions that occur during normal operation.

The small amount of CH₄ lost as ambient leakage measured here may not be economically important, but when extrapolated to a global scale the loss of 0.19% of gas production (the production-weighted average loss) from offshore oil and gas production corresponds to a global emission of 0.8 Tg CH₄ yr⁻¹ (IEA, 2018). Currently, the Oil and Gas Climate Initiative (OGCI) estimates the global CH₄ emission from the oil and gas sector to be 1.6 Tg CH₄ yr⁻¹, based on the OGCI's own estimate that 0.32% of CH₄ extracted is lost (OGCI, 2018). This estimate represents data from 13 of the largest oil and gas producers and accounts for up-stream CH₄ emissions from flaring, venting and offshore oil loading for all operated gas and oil assets. **If a global CH₄ emission from ambient leakage of 0.19% estimated by this study (0.8 Tg CH₄ yr⁻¹) is added to the current global estimate from flaring, venting and offshore oil loading (1.6 Tg CH₄ yr⁻¹) the total CH₄ emission from offshore oil and gas production would increase significantly. It should be noted that the value of 0.19% is based on a very small sample size using a method that comes with significant uncertainty. Moreover, the median value of this study (6.8 g s⁻¹) is much smaller than the regional median emission estimate of 99 g s⁻¹ for the Malay Peninsula and 15 g s⁻¹ for Borneo (Nara et al., 2015), which suggests that the ambient leakage rate may be lower in the North Sea than other regions of the world.** This study does highlight the shortcomings of using emission factors which rely on *a-priori* knowledge of the source, in contrast with direct measurements that account for all emissions and better estimate total emissions. In conclusion, we suggest that additional measurements of offshore oil and gas production platform operations (e.g. Saudi Arabia, Brazil, Mexico, Norway and the United States) be conducted to better inform leakage estimates and that these measurements be used to improve the UK and global CH₄ emission inventories.

Author contributions

J. Staunton-Sykes, S. N. Riddick and N. R. P. Harris designed the experiment, S. N. Riddick, J. Staunton-Sykes, G. Allen, J. Pitt and G.L. Forster prepared equipment and calibrated the instruments, S. N. Riddick, J. Staunton-Sykes and G.L. Forster carried out the measurements and J. Staunton-Sykes, A. J. Manning, D. Lowry, and E. G. Nisbet provided analysis. D. L. Mauzerall and M. Celia were the project leaders and provided scientific oversight and guidance throughout the planning, implementation, collection, and data analysis processes. S. N. Riddick and D. L. Mauzerall wrote the paper with help from M. Celia, M. Kang and N. R. P. Harris and with contributions from all co-authors.

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