We thank reviewer #1 for their careful assessment and for recommending the manuscript for publication in ACP. We have addressed all comments as detailed below (our responses are indicated in blue font).

2. General comments

2.1 Structure and presentation

We acknowledge that the readability of the manuscript can be improved by introducing sub-sections and further sub-divisions to existing paragraphs.

We have done this as detailed below:

Section 3.2.3: two new sub-sections on *"Temporal similarities"* and *"Comparison of flux spatial variability at the elevated and roof-top sites"*.

Section 3.3: one new sub-section added "3.3.1 Dependence of flux magnitude and diurnal patterns on wind sector".

Section 3.4: two new sub-sections created "3.4.1 Seasonal controls of fluxes of carbon monoxide and carbon dioxide" and "3.4.2 Seasonal controls of methane emissions".

Conclusion: we agree that the bullet point structure did give the impression of a draft document. This has now been re-structured into paragraph form.

2.2 Uncertainty quantification

The caption of Fig. 8 has been expanded and now includes a definition of the uncertainty ranges. In addition, this definition has also been added in section 3.5 for clarity.

As requested, confidence intervals have been added to Fig. 4 and absolute flux values are presented for CO₂ and CH₄ instead of the normalised ones originally used.

The estimated uncertainties on annual fluxes presented in Fig. 8 are of the order of 16% of the mean flux value for both CO_2 and CH_4 which is > 3 times the overall discrepancy between fluxes of CO_2 obtained from the open- and closed-path analysers, ca. 3 times the measured inter-annual variability in CO_2 fluxes and 4 times the inter-annual variability in CH_4 fluxes. There are relatively few uncertainty analyses for eddy-covariance fluxes measured on tall towers; instead, the standard deviation of the mean is commonly used to quantify inter-seasonal or inter-annual variability. Rigorous uncertainty analyses have been carried out for some tall tower sites but, to our knowledge, these case studies were only for homogeneous terrain (e.g. Post et al., 2015; Vickers et al., 2010) and it is thus difficult to put our uncertainty estimates from a highly heterogeneous environment into context.

2.3 Use of supplementary material

We agree with the reviewer that the scientific discussion should not occur outside of the main article. The manuscript preparation guidelines for ACP state that "The supplement **shall contain only complementary information** but no scientific interpretations or findings/messages that would go beyond the contents of the manuscript". We believe that our use of the supplement fulfils this criterion: the manuscript can be read and understood without the figures in the supplement but we feel that the additional information provided might be of interest to the readers.

2.4 Wind sector analysis

- We agree that a land-use map or satellite image would aid interpretation of the data presented in the manuscript. A satellite image indicating the two measurement sites discussed in this paper has been added to the supplementary material (Fig. S1).
- The wind sectors were defined so as to strike a reasonable balance between temporal and spatial coverages; more explicitly, the 45-degree wind sectors were preferred due to the high measurement height and the substantial seasonal variability in footprint which would result in averaging over very large spatial domains, especially in autumn and winter. We also found that the linear correlations between fluxes measured at the BT tower and KCL (Fig. 5) were weaker for broader wind sectors (we compared 45 degree and 90 degree wind sectors). Based on these considerations, we opted for the 45 degree definition.

3. Technical and specific comments

- Acronyms used before being defined: this has been addressed.
- Closing bracket should not be followed by opening bracket: we could not find any specific editorial guidelines regarding this point but we agree that this formatting is clumsy. We have made the relevant changes throughout the manuscript.
- Page 2, l. 14-15: We agree that this is a long list. We selected 5 multi-year studies and removed 3 references which are cited elsewhere in the manuscript.
- Page 4, line 28: the errors are directly averaged as stated.
- Page 6, section 3.2.1: there was nothing special about the data presented for that day. We selected relatively short averaging periods (e.g. 8-12 hours) to minimise potential effects from e.g. varying atmospheric stability, turbulence, relative humidity. We did not test for differences in co-spectral characteristics which could potentially arise over longer averaging periods; instead corrections for high-frequency attenuation were applied on a point-by-point basis for each half-hourly averaging period. We did however compare two contrasting relative humidity regimes, as mentioned in the manuscript, but found no compelling differences in the respective co-spectra.
- Page 6, line 23-32: the argument was expanded to "An explicit treatment of the storage term based on a gradient approach where concentrations and wind speeds are recorded at multiple heights below the EC measurement point could help probe the low u* regime. Such additional measurements were however not available for the BT tower site and we therefore speculate that the observation of venting after onset of turbulence, when the boundary layer grows, would capture at least some if not most of the material stored below the measurement height."
- Page 9, lines 9-10: We thank the reviewer for pointing out this ambiguous wording. The sentence has been changed to "Assuming that constant pressure is maintained throughout the gas distribution network, the leak rate should be constant through the day".
- Page 11, lines 3-18: we feel that since the temporal dynamics at diurnal and seasonal scales are attributed to distinct mechanisms, the discussion of the results benefits from keeping them separate. The readability has however already been improved by the introduction of new sub-sections.
- Page 12, lines 16-17: This is an important point to clarify and we thank the reviewer for bringing it up. We have added the following sentences to clarify the population argument: "Socio-economic temporal dynamics, such as a significant daytime influx of commuters into a business district (e.g. the City of London financial district which is located 3-4 km S-SE of the BT tower), might contribute substantially to CH₄ emissions (e.g. from sewage, natural gas); in addition, the measured CH₄ emissions from such business areas might bear no correlation

with the actual resident population reported here (source: London Datastore, Greater London Authority, 2016) which can be considerably smaller than the commuting workforce".

References

Post, H., H. J. H. Franssen, et al. (2015). "Uncertainty analysis of eddy covariance CO2 flux measurements for different EC tower distances using an extended two-tower approach." <u>Biogeosciences</u> **12**(4): 1205-1221.

Vickers, D., M. Gockede, et al. (2010). "Uncertainty estimates for 1-h averaged turbulence fluxes of carbon dioxide, latent heat and sensible heat." <u>Tellus Series B-Chemical and Physical Meteorology</u> **62**(2): 87-99.

We thank the reviewer for the positive review of our manuscript and for recommending its publication in ACP. We have addressed all comments as detailed below (our responses are indicated in blue font).

1. General comments

- Details of the KCL methane measurements using the QCL analyser: Section 2.2 on instrumentation has been further sub-divided into section "2.2.1 BT tower site" and "2.2.2 King's College London site" and the following description of the QCL eddy-covariance system at KCL has been added: "The QCL measured methane, nitrous oxide (N₂O) and water vapour simultaneously and at 10 Hz. The instrument was housed in an air-conditioned cabinet to minimise temperature fluctuations. Air was sampled ca. 20 cm below the anemometer head at 15 lpm through a 25 m long Teflon tube with outer diameter 1.27 cm (1/2"). The data was logged using an in-house LabView program and processed offline as outlined in section 2.3."
- The KCL CH₄ fluxes were processed in the same way as the BT tower fluxes. A sentence to this effect has been added to the instrumentation description (see new section 2.2.2 discussed above). The CO₂ fluxes from the KCL site were processed separately as detailed in Kotthaus and Grimmond (2014). The sentence on Page 4, line 5 was disambiguated by changing "*EC_kCL*" it to "*Fluxes of CO₂ from the KCL site*".
- Number of data points available at each site: The number of flux data points available for analysis is indeed a very important piece of information. We have added histograms detailing the number of half-hourly data points as a function of time of day and month of the year for each gas in the Supplementary material (Fig. S11-S14).
- We thank the reviewer for pointing out that the data periods used in Fig. 4 have not been defined in the caption. This has now been rectified in the figure caption and in the body of the manuscript: "The near-synchronous rise in CO₂ and CH₄ fluxes observed in summer (summer defined as the months (JJA) in the data period 15/09/2011 31/12/2013 for F_{CO2} and the entire period 19/08 01/10/2015 for F_{CH4}) at the two measurement sites (BT and KCL) at different heights is consistent with an earlier onset of turbulent mixing (Fig. 4 b, c)." Please also note that we revised the statement about the summertime fluxes of CH₄ at the low and high sites; after replotting F_{CH4} on absolute scales (Fig.4) and inserting confidence intervals as per reviewer 1's suggestions we deemed that the differences between the two measurement heights in terms of the onset of the early morning increase in CH₄ emissions were not compelling. This has no impact on the other findings or the discussion of the results.
- Correction of high frequency attenuation: the fluxes were only corrected once for highfrequency attenuation and we apologise for this unfortunate misunderstanding. The results section was heavily edited to eliminate unnecessary repetitions. In an effort to improve readability, the sections pertaining to high frequency attenuation and the comparison between open-path and closed-path systems were moved and merged into the materials and methods as suggested by reviewer #3. We trust that the new structure of the manuscript will prevent further misunderstandings regarding the corrections applied to the raw fluxes.
- Comparison of closed- and open-path analysers: we acknowledge that uncertainties can arise from both the closed-path and the open-path analyser. This has been clarified in the manuscript: "Increased scatter in F_{CO2_CP}, especially during low fluxes, could be due to uncertainties in determining the time-lag through maximisation of the covariance or also to uncertainties arising from the open-path analyser".
- Discussion of CO flux: we could not find any directly comparable study for CO which is why the emphasis was put on discussing its temporal trends and comparing our measurements with atmospheric emissions inventory data.

2. Minor comments

- Page 2, line 3: the sentence has been amended as suggested.
- Page 2, lines 8-9: we are not sure how to interpret this comment; the other two reviewers did not have concerns with that particular sentence which was hence left unchanged.
- Page 2, line 19: this has been addressed.
- Page 3, line 31: the acronym "EC" has now been defined at the first instance of use.
- Page 4, line 5: this instance of "EC_KCL" has already been replaced by "Fluxes of CO₂ from the KCL site" to address a comment from reviewer #1.
- Page 5, line 14: "30-minute" has been changed to "half-hourly" as suggested.
- Page 5, line 20: the correct terminology is "non-neutral stratification". This has been corrected in the manuscript.
- Page 6, lines 9-11: we thank the reviewer for spotting this typo. Instead of Co(wT), the text should read Co(wCO₂) and Co(wCH₄). This has been corrected.
- Page 9, lines 32-33: the 2015 summer measurement campaign for CH₄ lasted just under 2 months and we did not have enough data to compare all 8 wind sectors as we did for CO₂ for which we have > 2 years of simultaneous data at the two sites. We therefore cannot elucidate this question.
- Figure 4: As also requested by reviewer #1, confidence intervals have been added to Fig. 4 and absolute flux values are presented for CO₂ and CH₄ instead of the normalised ones originally used.
- Figure 5: 1) the data presented here are not daily means but diurnal averages (i.e. 24 bulkaveraged hourly values). The caption has been expanded for clarification. 2) using 30-minute flux data instead of the diurnal profiles generates much more scatter as can be expected. However, the wind sector segregated correlations between the 2 measurement heights remain broadly linear. 3) A satellite image with the locations of the BT tower and KCL has been added into the supplement. This image provides a visual aid for the readers from which sectors of potentially overlapping footprints can be deduced.
- Figure 7: We have added the following text to section 3.4 "The lowest emissions of CO were observed in April but this is thought to be an artefact caused by relatively low temporal and spatial coverage for that month resulting from instrument downtime. Whilst not used in the discussion that follows, the April data point is included in Fig. 7 c and f for consistency".
- Page 16, line 11 and page 18, line 7: spaces added.

References

Kotthaus, S., and Grimmond, C. S. B.: Energy exchange in a dense urban environment – part II: Impact of spatial heterogeneity of the surface, Urban Climate, 10, Part 2, 281-307, http://dx.doi.org/10.1016/j.uclim.2013.10.001, 2014.

We wish to thank the reviewer for their thorough and constructive assessment of our manuscript and for recommending its publication in ACP. We have addressed all comments as detailed below (our responses are indicated in blue font).

1. General comments

We acknowledge that the readability of the manuscript will benefit from editing and re-grouping sections in such a way as to avoid unnecessary repetitions. We have done this following the suggestions provided by the reviewer: in particular, the sections pertaining to high frequency attenuation and the comparison between open-path and closed-path systems were moved and merged into the materials and methods.

2. Specific comments

• Organisation of sections 2 and 3:

The sub-sections on corrections for high frequency attenuation and the comparison between the open- and closed-path sensors have been moved from section 3 to section 2. This improves readability, reduces repetition and also addresses the comment of reviewer #2 who found the original presentation of the flux correction procedure confusing.

• Applicability of eddy-covariance: "With regards to the methodology, I am concerned that the low data rate, heterogeneous surface, and tall measurement height violate the requirements of sound eddy-covariance measurements. The comparisons with the closed-path sensors elsewhere on the BT tower and at King's College offer nice checks and allay some of my concern."

Eddy-covariance measurements are routinely conducted atop tall towers in natural and urban environments (e.g. Davis, Global Change Biology, 2003; Haszpra, Agriculture and Forest Meteorology, 2005; Liu, Atmospheric Chemistry and Physics, 2012) and previous work at the BT tower site should reassure the reviewer of the soundness of the experimental approach: for example, Wood et al. (Boundary Layer Meteorology, 2010) showed that the sensor height at the BT tower is above the mean roughness sublayer depth and that the similarity theory, which is traditionally applied to flat, homogeneous rural terrain, is also applicable at our urban site. The emissions measured at the BT tower represent processes at the local scale (see e.g. Grimmond and Oke, Journal of Applied Meteorology, 1999) as exemplified by the observation of summertime photosynthetic uptake of CO_2 from the wind sectors entraining large parks (Helfter et al., Atmospheric Chemistry and Physics, 2011). The good agreement in annual CO_2 emissions between the atmospheric emissions inventory and the direct measurements at the BT tower in our current and previous (Helfter et al., Atmospheric Chemistry and Physics, 2011) studies demonstrates that a) the use of eddy-covariance at this tall tower site is justified, and b) the effects of the low sampling rate on measured fluxes are well corrected.

• "It would be nice to have a better way to check for vertical storage errors":

We agree with the reviewer on this point but in the absence of direct measurements of the storage term at our site we must rely on the indirect assumptions discussed in the manuscript. We are planning on building a gradient system at the BT tower site which will enable us to calculate the storage term in future.

• "I am very concerned that the header of Table 1 suggests that sonic anemometer data was missing for 6 months of the analysis period":

The ultrasonic anemometer was indeed out of action for ca. 6 months but thanks to the otherwise long time series we do have temporal "replicates" for the missing months.

3. Technical comments

• Abstract – The second half has too much detail, especially when you start to talk about the different wind sectors. The abstract should also answer the question of why this work is important. The last sentence currently is not an adequate answer to that question.

We acknowledge the weakness of the abstract which has been revised as follows (new text in italics):

"We report on more than three years of measurements of fluxes of methane (CH₄), carbon monoxide (CO) and carbon dioxide (CO₂) taken by eddy-covariance in central London, UK. Inter-annual variability in the period 2012-2014 ranged from 36.3 to 40.7 ktons km⁻² y⁻¹ for CO₂, and from 69 to 75 tons km⁻² y^{-1} for CH₄. Mean annual emissions of CO₂ (39.1 ± 2.4 ktons km⁻² y⁻¹) and CO (89 ± 16 tons km⁻² y⁻¹) were consistent (within 1% and 5% respectively) with values from the London Atmospheric Emissions Inventory, but measured CH_4 (72 ± 3 tons km⁻² y⁻¹) was over two-fold larger than the inventory value. Seasonal variability was large for CO with a winter to summer reduction of 69%. Monthly fluxes of CO were strongly anti-correlated with mean air temperature, and the winter emissions accounted for 45% of the annual budget. The winter increment in CO emissions was attributed mainly to vehicle cold starts and reduced fuel combustion efficiency. CO₂ fluxes were 33% higher in winter and anticorrelated with mean air temperature, albeit to a lesser extent than for CO. This was attributed to an increased demand for natural gas for heating during the winter. The seasonality in CH₄ fluxes was moderate (21% larger in winter) and the linear anti-correlation with air temperature was spatiallyvariable. Differences in resident population within the flux footprint explained up to 90% of the spatial variability of the annual CO₂ fluxes and up to 99% for CH₄. This suggests a significant influence of anthropogenic sources in the overall emissions budget of these two greenhouse gases. Furthermore, we suggest that biogenic sources of CH₄, such as waste water which is unaccounted for by the atmospheric emissions inventories, make a substantial contribution to the overall budget and that commuting dynamics in and out of central business districts could explain some of the spatial and temporal variability of the emissions. To our knowledge, this study is unique given the length of the datasets presented, especially for CO and CH₄ fluxes. This study offers an independent verification of "bottom-up" emissions inventories and demonstrates that the urban sources of CO and CO_2 are well characterised in London. This is however not the case for CH_4 emissions which are heavilyunderestimated by the inventory approach. This opens up opportunities in the UK and abroad to identify and quantify the "missing" sources of urban methane, revise the methodologies of the atmospheric inventories and devise emission reduction strategies for this potent greenhouse gas."

 Pg 1 – In the context of the motivation given for the study, I wonder whether eddy-flux is the really right tool for emissions verification. In natural systems, where it has been most widely applied, there is the problem of the representivity of individual towers and up-scaling across heterogeneous ecosystems. In urban environments, this problem is even more acute. Eddyflux in cities seems like it would be more useful for investigating mechanisms and drivers of emissions.

We have already provided an answer to question in section "2. Specific comments" but we wish to reiterate that a) the applicability of the eddy-covariance method for tall tower urban studies has been established by us and others in London and elsewhere, and that b) spatial and temporal emission mechanisms and drivers are discussed in this paper.

• Pg 1, In 10 – This reference of 70% of emissions is from cities is often inappropriately used to motivate urban studies. It should say 70% is attributable to cities because the number includes not only emissions that emanate directly from cities, but also upstream emissions, such as for power production, that occur outside of cities. Therefore, atmospheric measurements located in cities would not necessarily be sensitive to 70% of anthropogenic emissions.

We agree with that statement but we are merely quoting the literature on that point.

• Pg 1, ln 11 – "verify GHG emissions" could be changed to "evaluate reported GHG emissions"

We changed the wording as suggested.

• Pg 3, In 6 – You give the mean building height in the vicinity of the measurement, but what is the maximum building height? This would seem important to ensure that the air flow is not impeded around the sensor.

The references cited provide more details about the local topography and the turbulence and air flow characteristics observed at the BT tower. We therefore hope that these references are sufficient.

• Pg 3, In 17 – How do you know that weekly calibration was adequate? Your standard gases were mixtures in nitrogen, but the CRDS requires natural air standards. See Nara et al. 2012 (www.atmos-meas-tech.net/5/2689/2012/amt-5-2689-2012.html).

We are aware of that publication and have evaluated the performance of our Picarro CRDS in this light. The temporal drift of the instrument over a one week period is almost negligible and we are therefore reassured that weekly calibrations are sufficient. Furthermore, we have used natural compressed air as calibration gases since late 2014. The "matrix" effect reported by Nara is probably critical in the case of high-precision concentration measurements but we did not observe any significant impact on our flux measurements which are derived from relative changes in concentrations.

• Pg 5, ln 21 – A semicolon is not necessary. A period will suffice.

Agreed.

• Pg 5, ln 25 – Is the seasonal difference in the length-scale of the footprint, mainly a function of wind-speed?

Atmospheric stability and local turbulence also contribute to the footprint.

• Pg 5, ln 27 – It would be nice to include a map to help the reader visualize the nearby landscape.

This point was also raised by reviewer #2 and a satellite image centred on the BT tower was included as Fig. S1 of the supplement.

• Pg 6, In 19 – Not just increased scatter but the residuals look like they are biased high. Could this affect the mean results?

This is an interesting point which relies on the assumption that the open-path analyser is a perfect reference. The open-path analyser is not calibrated and the two sets of eddy-covariance data were analysed separately using different software.

• Pg 7, ln 2 – This would also result in the wrong time of day being attributed to the flux.

That is correct but this transient effect would be smoothed out a longer averaging intervals (e.g. monthly and annual).

• Pg 8, In 7 – How long do the periods need to be for this assumption to hold? This requirement to analyze long periods only seems counter to the discussion of hour-to-hour differences in the previous paragraph.

One full 24-hour cycle would in theory be sufficient to cancel out the storage-venting effect; this transient nocturnal storage term would be very important to account for if one were to integrate the hourly fluxes in order to derive daily budgets or attempt to partition emissions into daytime and night time contributions. This doesn't preclude discussing hour-to-hour difference but it must be remembered that the early morning peak in emissions sometimes observed at the BT tower can be a result of the growth of the boundary layer.

• Pg 9, In 8 – insert "natural" gas.

Done.

• Pg 9, ln 2 – "pollutants" could be changed to "gases".

Done.

• Pg 9, In 20 – Suggest deleting the statement about the gas supply pressure since it is highly speculative and you have zero information about actual supply line pressures. All that you can really say is that methane fluxes (natural gas or other) appear to vary diurnally.

We acknowledge that we do not have quantitative information regarding the pressurisation of the natural gas distribution network in London. However, this does not preclude proposing a mechanism to explain the observed diurnal variation in methane fluxes. The London atmospheric emissions inventory sets natural gas as the main contributor of atmospheric methane and we use these data as a starting point to our discussion. For these reasons, we have decided to leave that statement in the revised document.

• Pg 9, In 22 – Insert paragraph break at "Segregating"

This has been done and a new numbered section (3.3.1 Dependence of flux magnitude and diurnal patterns on wind sector) was introduced in order to improve the readability of the document.

• Pg 10, ln 11 – Could you actually predict the seasonal difference in CO emissions using the known temperature-dependence of vehicle CO emissions, temperature and vehicle-count data, or is this already done in the inventory model?

This is essentially how inventory data is obtained with the exception that the reported values are annual ones. The London Atmospheric Emissions Inventory does supply a breakdown of the terms which contribute to the total annual budget with cold starts and seasonal effect being accounted for.

• Pg 10, In 22 – Why the difference between your result and the previous London study? It might be worth briefly summarizing previous results for London in the introduction.

The difference between the present and previous studies falls within the uncertainty range of the annual budgets. Previous results for London are already summarised in Table 1 and we would rather avoid increasing the length of the paper by duplicating these numbers.

• Pg 11, ln 11 – Seasonal difference in natural gas emission in Boston were not significant.

We thank the reviewer for specifying this. The sentence was amended accordingly.

• Pg 11, ln 13 – "Although individually small" – incomplete sentence.

The sentence "Although individually small, passive fugitive post-meter emissions (i.e. in homes or work place) can make a non-negligible cumulative contribution at the city scale (Wennberg et al., 2012)." does seem complete.

• Pg 11, In 15 – Post-meter emissions could vary seasonally if they are correlated with appliance use.

The fugitive emissions discussed here occur constantly, irrespective of appliance use.

• Pg 13, ln 13 – Why are natural gas leaks any more likely as a source than sewage, for example, which could also be correlated with air temperature?

This was an oversight and sewage as a potential temperature-dependent source of methane has been included in the sentence.

• Pg 13, ln 15 – Change "the easiest to get right" to "most accurately represented".

Done. Thank you for suggesting this change of wording.

• Table 1 – The "Aircraft measurements" footnote should be below the table.

This has been changed.

• Figure 1 – Use units of km instead of m so that plot is not so cluttered with zeros.

This has been addressed.

• Figure 4 – Could just label each sub-panel so hatched lines are not needed.

The sub-panels have been individually labelled and solid lines are used throughout.

• Figure 5 – I do not know what "diurnal average" means. What does each point on the plot represent?

We acknowledge that the wording was ambiguous; "diurnal averages" have been replaced with "average diurnal profiles of CO₂ fluxes".

• Figure 6 – The y-axes on several panels have unnecessarily large ranges.

In our opinion, the ranges of the y-axes in Fig. 6 g-i allow for a direct comparison of the weekday-toweekend decrease in emissions for the three gases. The ranges of the y-axes in Fig. 6 a-c were chosen to be consistent with panels d-f. For these reasons, we have decided not to change the ranges of the axes in Fig. 6.

References

Davis, K. J., et al. (2003). "The annual cycles of CO_2 and H_2O exchange over a northern mixed forest as observed from a very tall tower." <u>Global Change Biology</u> **9**(9): 1278-1293.

Grimmond, C. S. B. and T. R. Oke (1999). "Aerodynamic properties of urban areas derived, from analysis of surface form." Journal of Applied Meteorology **38**(9): 1262-1292.

Haszpra, L., et al. (2005). "Long-term tall tower carbon dioxide flux monitoring over an area of mixed vegetation." <u>Agricultural and Forest Meteorology</u> **132**(1-2): 58-77.

Helfter, C., et al. (2011). "Controls of carbon dioxide concentrations and fluxes above central London." <u>Atmospheric Chemistry and Physics</u> **11**(5): 1913-1928.

Liu, H. Z., et al. (2012). "Four-year (2006-2009) eddy covariance measurements of CO_2 flux over an urban area in Beijing." <u>Atmospheric Chemistry and Physics</u> **12**(17): 7881-7892.

Wood, C. R., et al. (2010). "Turbulent Flow at 190 m Height Above London During 2006-2008: A Climatology and the Applicability of Similarity Theory." <u>Boundary-Layer Meteorology</u> **137**(1): 77-96.

Spatial and temporal variability of urban fluxes of methane, carbon monoxide and carbon dioxide above London, UK.

3

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14 Abstract. We report on more than three years of measurements of fluxes of methane (CH_4), carbon monoxide (CO) and carbon 15 dioxide (CO₂) taken by eddy-covariance in central London, UK. Inter-annual variability in the period 2012-2014 ranged from 36.3 to 40.7 ktons km⁻² v⁻¹ for CO₂, and from 69 to 75 tons km⁻² v⁻¹ for CH₄. Mean annual emissions of CO₂ (39.1 \pm 2.4 ktons 16 17 km⁻² y⁻¹) and CO (89 \pm 16 tons km⁻² y⁻¹) were consistent (within 1% and 5% respectively) with values from the London Atmospheric Emissions Inventory, but measured CH₄ (72 ± 3 tons km⁻² y⁻¹) was over two-fold larger than the inventory value. 18 19 Seasonal variability was large for CO with a winter to summer reduction of 69%. Monthly fluxes of CO were strongly anti-20 correlated with mean air temperature, and the winter emissions accounted for 45% of the annual budget. The winter increment 21 in CO emissions was attributed mainly to vehicle cold starts and reduced fuel combustion efficiency. CO₂ fluxes were 33% 22 higher in winter and anti-correlated with mean air temperature, albeit to a lesser extent than for CO. This was attributed to an 23 increased demand for natural gas for heating during the winter. Seasonality The seasonality in CH₄ fluxes was moderate (21% 24 larger in winter) and the linear anti-correlation with air temperature was only statistically significant for certain wind sectors 25 (N, NE, E and W), which was also the case for CO_2 was spatially-variable. Differences in resident population within the flux footprint explained ca.up to 90% of the spatial variability by wind direction in of the annual CO₂ fluxes and up to 99% for CH₄ 26 27 (wind sectors excluded from linear regressions: S for CO₂; S, SE and E for CH₄).-This suggests a significant influence of 28 anthropogenic sources in the overall emissions budget of these two greenhouse gases. Furthermore, we suggest that biogenic 29 sources of CH₄, such as waste water which is unaccounted for by the atmospheric emissions inventories, make a substantial 30 contribution to the overall budget and that commuting dynamics in and out of central business districts could explain some of the spatial and temporal variability of the emissions. Seasonality and proportionality of emissions with respect to population 31 32 in the outlying wind sectors (S, SE and E) might be masked by constant sources of CO₂ and CH₄, perhaps of industrial or 33 biogenic origin. To our knowledge, this study is unique given the long term, continuous ength of the datasets presented,

1 especially for CO of urbanand CH4 fluxes-analysed. This study offers an independent verification of "bottom-up" emissions

2 inventories and demonstrates that the urban sources of CO and CO₂ are well characterised in London. This is however not the

3 case for CH₄ emissions which are heavily-underestimated by the inventory approach. This opens up opportunities in the UK

4 and abroad to identify and quantify the "missing" sources of urban methane, revise the methodologies of the atmospheric

- 5 inventories and devise emission reduction strategies for this potent greenhouse gas.
- 6

7 1 Introduction

8 The use of eddy-covariance for the measurement of turbulent fluxes of heat and mass has grown steadily over the past three 9 decades; eurrently recently, there are were > 400 active sites worldwide (Baldocchi, 2008) spanning six continents. The vast 10 majority of existing sites were established to measure biosphere-atmosphere exchanges of carbon dioxide (CO₂) and heat-and carbon dioxide (CO₂) (Baldocchi et al., 2001). Due to recent technological advances, i.e. the development of new fast response 11 12 analysers, measurements of eddy-covariance fluxes of other trace gases such as methane (CH_4) and nitrous oxide (N_2O) are 13 gradually being introduced (Crosson, 2008; Fiddler et al., 2009; Peltola et al., 2014). With the negotiation of international 14 agreements to greatly reduce greenhouse gas (GHG) emissions by the end of the 21st century, there is an ever increasing need 15 to verify emissions through independent monitoring approaches. Despite 54% of the worldwide population currently living in cities, a figure which could rise to 66% by 2050 (United Nations, 2014), and urban CO₂ emissions estimated to represent 70% 16 17 of the global budget (International Energy Agency, 2012) there are comparatively few urban studies to evaluate reported GHG 18 emissionsverify GHG emissions. At the time of writing, 61 urban flux towers were listed in the FLUXNET Urban Flux 19 Network database, of which 40 were located in temperate areas (Grimmond and Christen, 2012). At present, most published 20 urban studies have focused on CO_2 at time scales ranging from a few months to a few years (e.g. Christen et al., 2011; Helfter 21 et al., 2011; Pawlak et al., 2011; Jarvi et al., 2012; Liu et al., 2012) (for recent publications and reviews see for example 22 Christen, 2014; Christen et al., 2011; Helfter et al., 2011; Jarvi et al., 2012; Liu et al., 2012; Pawlak et al., 2011; Velasco and 23 Roth, 2010; Ward et al., 2015). Methane, a potent GHG with a global warming potential 28 times larger than that of CO_2 at 24 the 100-year horizon (IPCC, 2013), is receiving increasing attention. Whilst CO₂ emissions are very closely linked to fuel 25 consumption, for which robust statistics can be obtained (at least at country level), CH₄ originates from a much larger range 26 of sources with complex controls. CH₄ emissions are commonly estimated in "bottom-up" inventories and at the national scale (e.g. for IPCC reporting), but also at the urban scale (e.g. London Atmospheric Emissions Inventory (LAEI) in the UK and the 27 28 California Air Resources Board (CARB) in the USA). A variety of techniques have recently been applied to provide 29 independent top-down estimates of urban CH₄ emissions. These include ground-based mass balance approaches (McKain et 30 al., 2015), airborne observations (O'Shea et al., 2014; Cambaliza et al., 2015), Fourier Transform Spectrometry (FTS) (Wunch 31 et al., 2009), isotopic source apportionment studies (e.g. Lowry et al., 2001; Zazzeri et al., 2015) and eddy-covariance (Gioli 32 et al., 2012; Pawlak and Fortuniak, 2016).

We report on over three years of continuous measurements of fluxes of methane, carbon monoxide and carbon dioxide in the heart of London, UK, the largest European city. This is, to our knowledge, the longest continuous urban record of direct CH₄ emission flux measurements. This paper investigates the temporal and spatial emission dynamics of the three pollutants and compares annual budgets with the bottom-up emissions inventory estimates.

5

6 2 Materials and methods

7 2.1 Site description

8 Fluxes of carbon monoxide (CO), carbon dioxide (CO₂) and methane (CH₄) were measured by eddy-covariance (EC) from the 9 rooftop of a 190 m telecommunication tower (BT tower; located at 51° 31' 17.4''N, 0° 8' 20.04''W) in central London, UK. 10 The measurements, which are ongoing at the time of writing, began in September 2011. The period September 2011 to 11 December 2014 is analysed here. The mean building height in a radius of ca. 10 km from the tower is 8.8 m ± 3.0 m and 12 typically 5.6 m ± 1.8 m for suburban areas (Wood et al., 2010; Evans, 2009). The Greater London area, which extends ca. 20 13 km in all directions from the BT tower, has a population of 8.6 million (Mayor of London Office, 2015) and population 14 densities in excess of 10⁴ inhabitants km⁻² in the central boroughs.

15 2.2 Instrumentation

16 <u>2.2.1 BT tower site</u>

17 The eddy-covariance system used at the BT tower consisted of a 3D ultrasonic anemometer (R3-50, Gill Instruments), a Picarro 18 cavity ringdown spectrometer (CRDS) model 1301-f for the measurement of CO₂, CH₄ and H₂O mole fractions and an 19 Aerolaser fast CO monitor model AL5002. The anemometer was mounted on top of a lattice tower located on the roof of the 20 BT tower giving an effective measurement height of 190 m above street level. The two gas analysers were located a few floors 21 below the roof, in an air conditioned room. Air was sampled from ca. 0.3 m below the anemometer head at 20-25 lpm using a 22 45 m long Teflon tube of OD 9.53 mm (3/8"). The Picarro CRDS was fitted with an in-house auto-calibration system and 23 calibrated weekly using two different mixtures of methane and carbon dioxide in nitrogen (above and below typical ambient 24 concentrations). The anemometer operated at 20 Hz, the CO analyser at 10 Hz and the Picarro CRDS, which was set to sample in 3-species mode, operated at 1 Hz. The data were captured by an in-house LabViewTM (National Instruments) data acquisition 25 26 program which also controlled the auto-calibration system and fluxes were processed offline by a custom LabView program. 27 Although the Picarro 1301-f has the capability to measure concentrations at 10 Hz, at this rate, this older instrument can only 28 measure two of the three compounds CO₂, CH₄ and H₂O. Because an internal H₂O measurement is required for accurate 29 corrections (e.g. Peltola et al., 2014), this would mean that in fast-response mode the instrument can only measure the flux of 30 CO_2 or CH_4 at any one time. Due to the high measurement height, it was found that a response time of 1 Hz was sufficient to 31 capture >70% of the flux (see below).

In addition to the closed-path system described above, an open-path infrared gas analyser (IRGA model Li7500, LI-COR Biosciences) measuring CO₂ and H₂O at 20 Hz was mounted next to the ultrasonic anemometer on the roof of the BT tower. Wind and IRGA data were saved to a datalogger and processed separately to the closed path dataBoth analysers used the same anemometer but data were processed independently with different eddy-covariance software packages. In what follows, subscripts "_CP" and "_OP" will respectively denote the closed-path and open-path eddy-covariance systems, and fluxes derived from them, located at the BT tower.

7

8 2.2.2 King's College London site

9 Fluxes of CO₂ measured by EC__{CP} (F_{CO2_CP}) were compared to fluxes measured at an eddy-covariance site at King's College 10 London (KCL; use of subscript "_{_KCL}" to identify this eddy-covariance system in what follows) Strand campus, about 2 km 11 south-east of the BT tower, where long-term EC measurements have been analysed to study energy exchanges (Kotthaus and 12 Grimmond, 2014a, b), carbon dioxide fluxes (Ward et al., 2015) and the F_{CO2} storage term in a dense urban environment 13 (Bjorkegren et al., 2015). Carbon dioxide fluxes are obtained from observations of an open-path Li7500 gas analyser and a 14 CSAT3 sonic anemometer (Campbell Scientific). KCL is within the flux footprint of the BT tower during south-easterly wind 15 directions, EC-_{KCL}Fluxes of CO₂ from the KCL site are-were processed as outlined by Kotthaus and Grimmond (2014a).

- 16 -For the time August September 2015, a methane sensor (Aerodyne Quantum Cascade Laser (QCL)) was added to the EC
- 17 system at KCL to also observe F_{CH4} . No carbon monoxide was measured at KCL. The EC_{KCL} system was operated at the top
- 18 of a tower situated on the roof of a large building resulting in a measurement height of 50 m above mean ground level in the
- 19 flux footprint (Ward et al., 2015), i.e. ca. 140 m lower than for EC_CP. Given that the KCL site is closer to the urban canopy,
- 20 the source area extends to several hundred metres, while the footprint of the BT tower is much larger, i.e. in the order of
- 21 kilometres. The QCL measured methane, nitrous oxide (N2O) and water vapour simultaneously and at 10 Hz. The instrument
- 22 was housed in an air-conditioned cabinet to minimise temperature fluctuations. Air was sampled ca. 20 cm below the
- anemometer head at 20 lpm through a 25 m long Teflon tube with outer diameter 1.27 cm (1/2^{''}). The data was logged using
- 24 an in-house LabView program and processed offline as outlined in section 2.3.

25 **2.3 Data processing and filtering**

- Half-hourly fluxes were calculated using standard eddy-covariance methodology extensively described elsewhere (e.g.
 Aubinet et al., 2000; Foken et al., 2004; Moncrieff et al., 2004).
- 28 2.3.1 High frequency attenuation
- 29 Normalised cospectra ('Co(x)') of wCO₂ and wCH₄ measured by the closed-path system were corrected to match those of wT
- 30 (where T is sonic temperature) to assess high frequency damping caused by the instrument's limited sampling rate (1 Hz),
- 31 internal instrument time response and the long inlet line (~ 45 m). Co(wT) followed the theoretical f^{-5/3} (where f denotes
- 32 frequency) slope for the inertial sub-range (Foken, 2008) over the entire frequency range (Fig. 2). In contrast, Co(wCO₂) and

1	$Co(wCH_4)$ diverged from the theoretical slope for frequencies > 0.1 Hz and followed profiles with slopes of the order of ~ f ⁻
2	$\frac{5/2}{2}$. Relative humidity did not have a significant influence on Co(w TCO₂) and Co(wCH ₄) for the two regimes tested (RH =
3	52% and RH = 80%, data not shown) which suggests that the dominant causes of signal attenuation for our system were the
4	sampling rate and the length of the inlet line. Typical corrections for high frequency attenuation ranged from 15%-30% and
5	based on the co-spectra presented in Fig. 2 it can be inferred that eddies of frequency < 0.1 Hz carried $> 70\%$ of the flux
6	measured at the 190 m above-street-level sampling height. The net flux loss resulting from high frequency attenuation was of
7	the order of 30% over the entire frequency range. Each half-hourly flux was corrected for high frequency attenuation as part
8	of the offline data processing procedure on a point per point basis.
9	
10	2.3.2 Quality control and filtering
11	Half-hourly means were excluded if any of the following quality assurance criteria were not fulfilled:
12	• The number of raw data points per nominal half hour was < 35000.
13	• The flow rate in the sampling line was less than 15 lpm (theoretical limit of the transitional phase between laminar
14	and turbulent flow for the sampling tube diameter used in this study).
15	• The number of spikes in u, v, w (components of the 3D wind vector measured by the ultrasonic anemometer) or any
16	of the trace gas mole fractions was > 360 (i.e. 1% threshold).
17	• Latent and sensible heat fluxes fell outside the -250 W m ⁻² to +800 W m ⁻² range.
18	• The level of turbulence was deemed insufficient for flux measurement (friction velocity, u_* , threshold of 0.2 m s ⁻¹).
19	This threshold was used for consistency with previous studies carried out at the BT site (Helfter et al., 2011; Langford
20	<u>et al., 2010).</u>
21	• The stationarity test which requires that the difference between the half-hourly flux and the fluxes obtained from 6 x
22	5 min averaging sub-intervals does not exceed 30% is satisfied (Foken and Wichura, 1996; Foken, 2004).
23	2.3.3 Comparison between closed-path and open-path systems
24	The performance of the closed-path greenhouse gas eddy-covariance system located on the 35 th floor of the BT tower (EC_CP)
24 25	The performance of the closed-path greenhouse gas eddy-covariance system located on the S5 floor of the B1 tower (EC_CP) was compared to that of the open-path IRGA located on the roof of the tower (EC_OP). EC_CP operated at 1 Hz, EC_OP at 20
23 26	Hz and.
20 27	After frequency correction, half-hourly CO_2 fluxes measured by an open-path Li7500 infrared gas analyser located on the roof
28	of the BT tower ($F_{CO2 OP}$) were strongly correlated to the fluxes obtained with the closed-path Picarro analyser ($F_{CO2 OP}$; Fig.
20 29	3). Increased scatter in F_{CO2_CP} , especially during low fluxes, could be due to uncertainties in determining the time-lag through
30	maximisation of the covariance or also to uncertainties arising from the open-path analyser. The slope of near unity indicates
	5

1	that the high frequency attenuation of the turbulent flow due to instrument response time, sampling flow rate and length of the
2	sampling line was adequately and systematically corrected for.
3	<u>F_{CO2_CP}</u> clearly varied with friction velocity (u_*) with maximum fluxes observed at u_* values around 0.8 m s ⁻¹ , strongly reduced
4	fluxes at low $u_{\underline{*}} < 0.3 \text{ m s}^{-1}$ and the indication of reduced values at very high values of $u_{\underline{*}}$ (Fig. S2). A similar $u_{\underline{*}}$ dependence
5	was found for the fluxes from the open-path gas analyser (not shown). Near-zero fluxes were recorded by both systems for u_*
6	<u>values $< 0.1 \text{ m s}^{-1}$. For CO₂ flux measurements over vegetation, this type of behaviour is usually attributed to a reduction in</u>
7	the transport to the measurement height, resulting in storage of CO ₂ below that height which may be subject to advection. In
8	the urban environment, this <i>u</i> * dependence could alternatively arise from an actual correlation between <i>u</i> * and surface emission.
9	Indeed, on average both u_* and traffic counts show a minimum at night. However, it is likely that a loss of coupling with street
10	level sources as a result of limited vertical transport occurred in situations of low turbulence. These situations often coincide
11	with stable night-time conditions during which the boundary layer height can approach that of the measurement height (Barlow
12	et al., 2015). In such conditions the measured flux would be an underestimate of the true surface emission due to change in
13	storage in the air column below the measurement height. An explicit treatment of the storage term based on a gradient approach
14	where concentrations and wind speeds are recorded at multiple heights below the EC measurement point could help probe the
15	low u _* regime. Such additional measurements were however not available for the BT tower site and we therefore speculate
16	that the observation of venting after onset of turbulence, when the boundary layer grows, would capture at least some if not
17	most of the material stored below the measurement height.
18	
19	
20	Half hourly means were excluded if any of the following quality assurance criteria were not fulfilled:
21	• The number of raw data points per nominal half hour was < 35000.
22	• The flow rate in the sampling line was less than 15 lpm (theoretical limit of the transitional phase between laminar
23	and turbulent flow for the sampling tube diameter used in this study).
24	• The number of spikes in u, v, w (components of the 3D wind vector measured by the ultrasonic anemometer) or any
25	of the trace gas mole fractions was > 360 (i.e. 1% threshold).
26	• Latent and sensible heat fluxes fell outside the 250 W m ⁻² to +800 W m ⁻² range.
27	• The level of turbulence was deemed insufficient for flux measurement (friction velocity, u_{\pm} , threshold of 0.2 m s ⁻¹).
28	This threshold was used for consistency with previous studies carried out at the BT site (Helfter et al., 2011; Langford
29	et al., 2010).

The stationarity test which requires that the difference between the half hourly flux and the fluxes obtained from 6 x
 <u>5 min averaging sub-intervals does not exceed 30% is satisfied (Foken and Wichura, 1996; Foken, 2004).</u>

3 2.4 Uncertainty analysis

1

2

Random measurement uncertainties were estimated for each half-hourly averaging period using the Finkelstein and Sims 4 5 (2001) method and subsequently averaged into monthly means. The upper bounds of the random uncertainties associated with 6 the annual emissions estimates were taken as the maximum monthly random uncertainty for each year and trace gas. 7 Unlike random uncertainties, which arise from instrument noise and representativeness of single-point measurements, 8 systematic errors can be minimised by careful data processing and correction. In particular, successive calibration events were 9 linearly interpolated over time, cancelling out errors due to calibration drifts assuming that the drift was linear over time. 10 A correction for high frequency attenuation of the 1 Hz eddy covariance signal was applied at the post processing stage and half-hourly fluxes of CO2 measured by the closed-path Picarro system were compared to fluxes obtained from a LICOR 7500 11 12 open path IRGA mounted near the ultrasonic anemometer on the rooftop of the BT tower. The magnitude of the correction for 13 high frequency attenuation was typically of the order of 15 20% of the flux; open and closed path fluxes agreed well post-14 correction and systematic uncertainties on processed half-hourly fluxes will hereafter be deemed negligible compared to 15 random uncertainties. 16 So far we have considered the error in the local flux. In addition, there is an uncertainty of how this local flux relates to the

16 So far we have considered the error in the local flux. In addition, there is an uncertainty of how this local flux relates to the 17 emission at the surface. The effects of advection and storage on the flux measurement are difficult to quantify in a 18 heterogeneous environment like a city. However, whilst individual 30 minutehalf-hourly flux values may be a poor 19 representation of the momentary emission, we expect the errors to reduce significantly when long-term averages are analysed. 20 The validity of this assumption is explored in more detail belowin what follows.

21 3 Results and discussion

22 **3.1 Flux footprint**

23 For consistency with a previous study (Helfter et al., 2011), the flux footprint for the BT tower measurement site was estimated 24 with the analytical model of Kormann and Meixner (2001) for non-neutral atmospheric stratification, under the simplifying 25 assumptions that fluxes of heat and momentum were homogenous across the footprint.; the The frequency of observation of 26 x_{90} , the distance from the tower where 90% of the measured fluxes originated from, is shown in Fig. 1 as a function of wind 27 direction and season for the measurement period 2011-2014. The spatial extent of the flux footprint was highly variable over 28 time with recurring seasonal patterns. Typically, 90% of the flux measured at the BT tower site originated from distances of 29 the order of a few km in spring and summer compared to several tens of km in winter. The flux footprint contains two large 30 parks in the SW (Hyde Park, surface area 142 ha) and NW (Regent's Park, surface area 197 ha), sub-urban residential areas in 1 the N, a mixture of heavily urbanised residential and commercial areas in the E and S and a section of the Thames river in the

2 SE.

3 3.2 Performance of the eddy-covariance system

4 The performance of the closed-path greenhouse gas eddy-covariance system located on the 35th floor of the BT tower (EC_CP)

5 was compared to that of the open path IRGA located on the roof of the tower (EC_OP). EC_CP operated at 1 Hz, EC_OP at 20

6 Hz and both analysers used the same anemometer data but were processed independently with different eddy covariance

7 software packages.

8 3.2.1 High frequency attenuation

Normalised cospectra (Co(x)) of wCO₂ and wCH₄ measured by the closed path system were corrected to match those of wT 9 (where T is sonic temperature) to assess high frequency damping caused by the instrument's limited sampling rate (1 Hz), 10 internal instrument time response and the long inlet line (~ 45 m). Co(wT) followed the theoretical f^{5/2} (where f denotes 11 frequency slope for the inertial sub-range (Foken, 2008) over the entire frequency range (Fig. 2). In contrast, Co(wCO₂) and 12 Co(wCH₄) diverged from the theoretical slope for frequencies > 0.1 Hz and followed profiles with slopes of the order of -f 13 ^{5/2}. Relative humidity did not have a significant influence on Co(wT) for the two regimes tested (RH = 52% and RH = 80%. 14 data not shown) which suggests that the dominant causes of signal attenuation for our system were the sampling rate and the 15 length of the inlet line. Typical corrections for high frequency attenuation ranged from 15%-30% and based on the co-spectra 16 presented in Fig. 2 it can be inferred that eddies of frequency < 0.1 Hz carried > 70% of the flux measured at the 190 m above-17 street-level sampling height. The net flux loss resulting from high frequency attenuation was of the order of 30% over the 18 19 entire frequency range. Each half-hourly flux was corrected for high frequency attenuation as part of the offline data processing procedure on a point per point basis. 20

21 3.2.2 Comparison between closed-path and open-path systems

After frequency correction, half hourly CO_2 fluxes measured by an open path Li7500 infrared gas analyser located on the roof of the BT tower (F_{CO2_OP}) were strongly correlated to the fluxes obtained with the closed path Picarro analyser (F_{CO2_CP}). (Fig. 3). Increased scatter in F_{CO2_CP} , especially during low fluxes, is likely due to uncertainties in determining the time lag through maximisation of the covariance. The slope of near unity indicates that the high frequency attenuation of the turbulent flow due to instrument response time, sampling flow rate and length of the sampling line was adequately and systematically corrected for.

- 28 F_{CO2_CP} clearly varied with friction velocity (μ_*) with maximum fluxes observed at μ_* values around 0.8 m s⁻¹, strongly reduced
- 29 fluxes at low $u_{\pm} < 0.3 \text{ m s}^{-1}$ and the indication of reduced values at very high values of u_{\pm} (Fig. S1). A similar u_{\pm} dependence
- 30 was found for the fluxes from the open path gas analyser (not shown). Near zero fluxes were recorded by both systems for u_*
- 31 values $< 0.1 \text{ m s}^{-1}$. For CO₂ flux measurements over vegetation, this type of behaviour is usually attributed to a reduction in

the transport to the measurement height, resulting in storage of CO_2 -below that height which may be subject to advection. In 1 2 the urban environment, this *u*^{*} dependence could alternatively arise from an actual correlation between *u*^{*} and surface emission. 3 Indeed, on average both μ_{*} and traffic counts show a minimum at night. However, it is likely that a loss of coupling with street 4 level sources as a result of limited vertical transport occurred in situations of low turbulence. These situations often coincide 5 with stable night time conditions during which the boundary layer height can approach that of the measurement height (Barlow et al., 2015). In such conditions the measured flux would be an underestimate of the true surface emission due to change in 6 7 storage in the air column below the measurement height. However, the observation of venting after onset of turbulence, when the boundary layer grows, would capture at least some if not most of the material stored below the measurement height. 8

9 3.2.3 Comparison with flux measurements at a lower height

10 3.2.1 Temporal similarities

F_{CO2_CP} and CO₂ fluxes observed at the KCL site (F_{CO2_KCL}) exhibited a high temporal correlation for diurnal patterns in both winter and summer (Fig. 4a, b; averaging period 15/09/2011 to 31/12/2013). Daily minima occurred at around 3:00 at both sites which is consistent with minimum traffic loads (Fig. 4f, g). Fluxes tended to increase from ca. 5:00-6:00 until late morning and declined steadily from ca. 18:00 at both sites, which is again in agreement with the declining traffic numbers in the evening. Methane fluxes exhibited similar temporal dynamics with the lowest emissions recorded during the night and a sharp rise between ca. 5:00 and 8:00. A gradual decrease in F_{CH4} was observed at both sites following the mid-morning maximum.

17 In winter, carbon dioxide fluxes started to increase slightly earlier (by about 30 min on average) at the KCL site. While this 18 time lag was not evident in the summer for F_{CO2} , methane fluxes started rising later at the elevated measurement point at BT 19 tower even in summer. Boundary layer growth in the morning transition period might explain some of the time delay observed 20 in the carbon fluxes. Mixing height (MH) estimates for several weeks in winter (6 Jan - 11 Feb 2012) and summer (23 July -21 17 Aug 2012) derived from Doppler LIDAR turbulence measurements (Bohnenstengel et al., 2015) at sites close to BT tower 22 (Fig. 4d, e) indicate that, on average, turbulent mixing extended above the BT tower measurement height of 190 m in both 23 seasons. However, mixing height exhibits great temporal variability depending on the synoptic background conditions; for 24 London it has been found that MH development depends primarily on the boundary layer winds and stability (Halios and 25 Barlow, 2016) so that these short-term climatology estimates might not be representative for the full period analysed for the 26 turbulent fluxes.

Growth of the convective layer was rapid in summer and a plateau was typically reached mid-morning which lasted until late afternoon. In agreement with the shorter day-length in winter, growth of the mixing height was slower, collapsing earlier in the evening after the mid-afternoon maximum. Daytime maximum mixing height was about 30% lower in winter compared to the summer. In both summer and winter, traffic counts rose during the morning transition period i.e. before the mixing layer started growing considerably (Fig. 4f, g); in the evening, traffic counts began decreasing after the mixing height had reduced in height. Given that the mean temporal evolution of carbon dioxide fluxes observed at both KCL and BT tower appeared to be closely linked to the profiles of road traffic, vehicle emissions apparently represent a significant control not only for the

local-scale observations at KCL (Ward et al., 2015) but also for fluxes at the elevated BT tower measurement point (Helfter et 1 2 al., 2011). The slight morning delay in wintertime F_{CO2} (Fig. 4a) observed at BT tower might be explained by the efficacy of 3 vertical turbulent transport between street level and the top of the BT tower which has been shown to depend on atmospheric 4 stability. The timescale of upward vertical turbulent transport was estimated to be of the order of 10 minutes for near-neutral 5 conditions, increasing to 20-50 minutes for stable conditions (Barlow et al., 2011). Low turbulence and prolonged periods of 6 stable atmospheric stratification (Fig. <u>\$2\$3</u>) could thus explain the 1-2 hour lag between the timing of the morning increase in 7 traffic counts and fluxes of CO_2 at the BT tower during the winter (Fig. 4a). This is consistent with the lag time observed for 8 profiles of potential temperature, and thus upward mixing, measured at the BT tower and a lower-level measurement site close 9 to the BT tower at 18 m a.g.l. (Barlow et al., 2015). The near-synchronous rise in CO₂ and CH₄, F_{CO2} fluxes observed in summer (summer defined as the months (JJA) in the data period 15/09/2011 - 31/12/2013 for F_{CO2} and the entire period 19/08 -10 01/10/2015 for F_{CH4}) at the two measurement sites (BT and KCL) at different heights is consistent with an earlier onset of 11 12 turbulent mixing g. This does however not explain the delay between F_{CH4} -measured at the two sites in summer 2015 (Fig. 4) 13 <u>b,</u>c).

14 Storage fluxes are difficult to quantify accurately in a heterogeneous environment like a city as this would require vertical 15 profile measurements below the measurement height at several locations within the flux footprint. The analysis presented here 16 therefore relies to some extent on the assumption that, over long periods, positive and negative storage fluxes cancel out and 17 that effects of advection on the stored quantity are negligible. This assumption is further supported by the very small storage 18 fluxes (< 2.5 % of the magnitude of the vertical fluxes) calculated at the KCL site (Bjorkegren et al., 2015), although these 19 would be somewhat larger for the higher measurement height at BT. While the turbulent fluxes observed at the BT tower and 20 KCL show close temporal alignment (Fig. 4a-c) their absolute values can differ considerably (e.g. KCL-to-BT ratios of peak 21 FCO₂ ranged from 1.5 in winter to 0.9 in summer; the summer ratio for FCH₄ was 1.5).

22

23 <u>3.2.2 Comparison of flux spatial variability at the elevated and roof-top sites</u>

24 Both sites are situated in central London where anthropogenic emissions are high due to the elevated density of people and 25 traffic (Ward et al., 2015). While the source area of the BT site includes central business district (CBD) areas with mostly 26 medium density midrise building structures, residential areas as well as large parks, the KCL footprint is dominated by CBD 27 structures with hardly any vegetation (Kotthaus and Grimmond, 2014b). Only the river Thames in its vicinity reduces 28 anthropogenic emission in some parts of the KCL footprint. To evaluate the response of F_{CO2} CP and F_{CO2} KCL to variations in 29 source area characteristics, the observations were grouped into eight sectors based on the wind direction measured at the BT 30 tower (Fig. 5). The carbon dioxide fluxes observed at the two sites are linearly correlated for all eight wind sectors but slope and goodness of fit vary. This is likely due to differences in flux footprints at the two measurement sites, including the extent 31 32 (a few km at the BT tower and a few hundred metres at KCL;- (Kotthaus and Grimmond, 2014b)) as well as emission source 33 density (a function of surface types). Near 1:1 agreement was found in the dominant SW wind sector (Fig. 5). For other wind 34 directions, differences in local-scale source area between the two EC sites become apparent: while a large green space

(Regent's Park) is located to the NW of BT tower, the surface seen by the KCL measurements is least urbanised towards the 1 2 S and SE of the site (river Thames; note that busy Waterloo bridge towards the SW of KCL acts as a very strong line source 3 of CO₂ keeping the fluxes relatively high from this wind direction). In response to the surface cover, F_{CO2} CP exceeds F_{CO2} KCL. 4 in the E, S and SE wind sectors by 20%, 50% and 70%, respectively, and is lower by 50%-70% in the N, NW and W sectors 5 with the poorest correlation for the NW sector. The smallest F_{CO2} CP fluxes were observed in the NW sector while F_{CO2} KCL. was highest in sectors NW and W where the particularly busy Aldwych junction is located (Kotthaus and Grimmond, 2014a). 6 7 KCL falls within the footprint of the BT tower site for SE wind direction, but clearly the BT tower measurement sees additional 8 sources due to the larger footprint. The focus was placed on discussing CO_2 fluxes in this section is because it is the only 9 compound for which we have a second long-term flux record at a lower measurement height. However, we assume that all the 10 conclusions (pertaining to e.g. spectral corrections, turbulent transport) in this section are also applicable to CH₄ and CO.

11 **3.3 Diurnal variability** of the measured fluxes

Fluxes-The fluxes of all three pollutants-gases (F_{CO} , F_{CO2} and F_{CH4}) exhibited well-defined diurnal cycles with minimum emissions during the night, typically from midnight until 5:00 GMT (Fig. 6a-c). Emissions increased sharply from 6:00 reaching a daytime maximum at around 12:00, and then declined steadily until early evening when a local maximum was observed at around 18:00-19:00. Mean F_{CH4} ranged from 5.7 to 11.0 kg km⁻² hour⁻¹ (maximum-to-minimum ratio of 1.9), F_{CO2} from 1867 to 6635 kg km⁻² hour⁻¹ (maximum-to-minimum ratio 3.5) and F_{CO} from 4.6 to 16.9 kg km⁻² hour⁻¹ (maximum-tominimum ratio 3.7), demonstrating that the relative dynamic range of F_{CH4} is less than that of the other compounds.

18 Urban CH₄ emissions from developed cities are thought to be dominated by leakage from the natural gas distribution network 19 and this is also reflected in the source sector breakdown of the LAEI. Assuming a-that constant pressure is maintained 20 throughout the gas distribution networkconstant gas supply pressure, the leak rate should be constant through the day. Thus, 21 Gioli et al. (2012) interpreted non-negligible diurnal cycles in F_{CH4} observed in Florence (Italy) to be caused by vertical 22 transport following the diurnal cycle of convective mixing. This argument does not seem relevant for the present study because 23 the diurnal variations in F_{CH4} measured at the BT tower were mirrored by strongly suppressed night-time CH₄ fluxes observed 24 at a much lower height at the KCL site (Fig. 4c), where the storage error has been demonstrated to be small for CO₂ (Bjorkegren 25 et al., 2015). Summer time fluxes of CH_4 measured at the 190 m height did lag slightly behind the fluxes observed at the 50 m 26 height, but this apparent delay could have been caused by differences in flux footprint between the sites (e.g. the source area 27 of the BT tower fluxes has a much higher fraction of vegetation cover than the KCL footprint) and the fact that the diurnal 28 profiles were obtained for a much shorter time period (August-September 2015). Instead, the similarity in F_{CH4} temporal 29 dynamics between the two sites supports the idea that the diurnal variations for that gas represent real variability in its source 30 strength rather than an artefact of atmospheric transport. This suggests that either the gas supply pressure in the distribution 31 network exhibits diurnal variations and/or that other CH₄ sources with temporal variations are more significant than estimated 32 by LAEI. This is further supported by F_{CH4} being smaller at the weekend than on weekdays (Fig. 6g).

1 3.3.1 Dependence of flux magnitude and diurnal patterns on wind sector

Segregating emissions by wind direction reveals heterogeneous source distributions at the BT tower site with different temporal 2 patterns (Fig. 6d-f) and source strengths (Fig. $\frac{83}{5}$ S4- $\frac{85}{5}$ S6). The lowest emissions (± standard error of the mean) for all three 3 4 pollutants were recorded for NW winds ($F_{CO} = 1.7 \pm 0.3$ kg km⁻² hour⁻¹, $F_{CO2} = 728 \pm 127$ kg km⁻² hour⁻¹, $F_{CH4} = 1.9 \pm 0.2$ kg km^{-2} hour⁻¹). The highest emissions of methane were found in the SE wind sector (17.8 ± 1.3 kg km⁻² hour⁻¹), in the S sector 5 for carbon dioxide (9020 \pm 515 kg km⁻² hour⁻¹) and in the E sector for carbon monoxide (25.4 \pm 3.9 kg km⁻² hour⁻¹). The 6 difference in emissions between wind sectors was however only statistically significant for the N and NW wind sectors. 7 8 Maxima of F_{CO} , F_{CO2} and F_{CH4} occurred on average at around 7:00-8:00 in the NW sector. Peak emissions for F_{CO2} and F_{CH4} in 9 the remaining wind sectors occurred typically between 9:00 and 12:00. The overall diurnal profile of F_{CO} was bimodal, except for NE and NW, with well-defined mid- to late-morning peaks (typically 9:00 to 12:00 GMT) followed by early evening peaks 10 11 (17:00 to 19:00). F_{CO} and F_{CO2} reached night time minima at around 3:00 in all wind sectors whereas F_{CH4} tended to plateau, 12 except in the SE where emissions tended to increase. The onset of an early morning increase in emissions (ca. 5:00-6:00 GMT) 13 was consistent for all wind directions for F_{CO2} and F_{CO} but it was less clearly defined for F_{CH4} . In addition to diurnal trends and dependency on wind sector, emissions of all three pollutants were found to be lower on weekends (Fig. 6g-i), with CH₄ again 14 showing the lowest variability (9% reduction on weekends compared to working days for F_{CH4} , 22% for F_{CO2} and 23% for 15 16 F_{CO}).

17 3.4 Seasonality of the measured fluxes trends

For the measurement period September 2011 to December 2014, F_{CH4}, F_{CO2} and F_{CO} exhibited marked seasonal cycles with 18 19 minimum emissions in summer (Fig. 7a-c). The lowest emissions of CO were observed in April but this is thought to be an artefact caused by relatively low temporal and spatial coverage for that month resulting from instrument downtime. Whilst not 20 21 used in the discussion that follows, the April data point is included in Fig. 7 c and f for consistency. For the months December-February, F_{CO2} and F_{CH4} were 4.1 ± 0.5 ktons km⁻² month⁻¹ and 7.4 ± 0.8 tons km⁻² month⁻¹, respectively, and decreased to 2.7 22 \pm 0.3 ktons km⁻² month⁻¹ (33% reduction) and 5.8 \pm 0.4 tons km⁻² month⁻¹ (21% reduction), respectively, in summer (June-23 24 August). The difference between winter and summertime emissions of carbon monoxide was 3-fold with 9.1 ± 2.5 tons km⁻² month⁻¹ in December-February and 2.9 ± 0.1 tons km⁻² month⁻¹ in June-July (due to instrument downtime, no data are available 25 26 for August).

27

28 3.4.1 Seasonal controls of fluxes of carbon monoxide and carbon dioxide

It is well established that emissions of CO from petrol cars are temperature dependent, e.g. increasing by a factor of 5-6 at ambient temperature 0 °C compared to 25 °C (Andrews, 2004) during the first 5-10 minutes following engine warm-up. The strong negative linear dependence of F_{CO} upon air temperature (Fig. 7f) could thus indicate that cold starts and reduced combustion efficiency played an important role during winter. Winter time (December-February) emissions of CO accounted for 45% of the annual budget for this pollutant which is consistent with LAEI (LAEI, 2010) estimates of the combined natural

gas and cold start contribution to annual CO emissions (total 32%, with 26% and 6% attributed to cold starts and natural gas 1 consumption, respectively). F_{CO2} was also correlated with air temperature (Fig. 7e; $R^2 = 0.59$), albeit to a lesser extent than 2 F_{CO} , which reflected the seasonal changes in domestic and commercial natural gas usage, but may also be influenced by 3 4 increased photosynthetic uptake by vegetation in the footprint during the warmer months. Anti-correlations between monthly 5 F_{CO2} and air temperature have been reported in other studies (e.g. Beijing, Liu et al., 2012; London, Ward et al., 2015). The gradient between F_{CO2} and air temperature observed in this study (-0.94 µmol m⁻² s⁻¹ °C⁻¹) falls between the values reported 6 for the London site (-1.95 μ mol m⁻² s⁻¹ °C⁻¹) and the Beijing site (-0.34 μ mol m⁻² s⁻¹ °C⁻¹). 7 8 The flux ratio of CO to CO_2 is of the order of 4 mmol mol⁻¹ in winter (excess CO due to cold starts and incomplete combustion)

9 and 2 mmol mol⁻¹ in summer despite only moderate seasonal variations in traffic loads (Fig. <u>S6S7</u>). Traffic loads at Marylebone 10 Road, one of the busiest arteries in central London located < 1 km north of the BT Tower, varied by less than 5% seasonally 11 in the period June 2012 to December 2014 (source Transport for London; personal communication). The seasonality of F_{CO2} 12 is hence likely controlled by changes in natural gas consumption and vegetation (Gioli et al., 2012; Helfter et al., 2011). This 13 is further supported by relatively constant ratios of F_{CH4} to F_{CO2} which suggests that seasonal variations in emissions were of 14 comparable magnitude for these two gases (Fig. <u>\$6</u>\$7). On average over the full three years of the study (2012-2014), 15 summertime F_{C02} were 30% lower than in winter (29% in 2012, 30% in 2013 and 2014). In comparison, during an earlier study 16 at the same site covering the year 2007, the winter to summer decrement was only 20% (Helfter et al., 2011).

17

18 3.4.2 Seasonal controls of methane emissions

19 Fluxes of CH₄ were 17% lower in summer than in winter (18%, 12% and 20% for 2012, 2013 and 2014 respectively) and the linear correlation of monthly averages with temperature was not statistically significant (Fig. 7d; $R^2 = 0.31$, p-value = 0.06). 20 21 In contrast, the winter to summer decrease was of the order of 63% in the city of Łódź, Poland (Pawlak and Fortuniak, 2016) and the dependence of F_{CH4} upon air temperature was statistically significant. The weaker correlation of F_{CH4} with air 22 23 temperature in London suggests that the total methane flux is due to a superposition of sources with constant and time-varying 24 emission rates, whereas in Florence (Italy) no significant seasonality in CH₄ emissions was observed (Gioli et al., 2012). They 25 related this to a constant pressure in the gas distribution network serving Florence. However, seasonality in both methane 26 concentrations and methane isotopic signature have been reported in the Greater London area (Lowry et al., 2001). The winter 27 time increase above background in CH₄ concentrations and the accompanying enrichment in δ^{13} C were consistent with North 28 Sea natural gas and attributed to losses of CH₄ from over-pressurised pipelines in response to (or anticipation of) an increase 29 in demand and to incomplete combustion upon boiler ignition. The seasonality of F_{CH4} in Łódź (Poland) was also attributed to 30 variations in natural gas usage (Pawlak and Fortuniak, 2016). Urban CH₄ emissions in Boston (USA) attributed to natural gas 31 use also displayed a modest, albeit not statistically significant, seasonality, with lower emissions during the summer (McKain 32 et al., 2015). An increase of total CH_4 emissions in summer could indicate temperature-sensitive biogenic sources played an 33 important role in Boston. Although individually small, passive fugitive post-meter emissions (i.e. in homes or work place) can 34 make a non-negligible cumulative contribution at the city scale (Wennberg et al., 2012). If post-meter emissions are constant in time, they would be part of the baseline CH₄ rather than being seasonally variable. Finally, methane emissions from liquefied
petroleum gas (LPG) vehicles, although small compared to natural gas emissions, exhibit a positive dependence upon
temperature (Nam et al., 2004) and are expected to also contribute to the seasonality and diurnal variation of the total urban
CH₄ fluxes.

5 3.5 Annual budgets of methane, carbon monoxide and carbon dioxide emissions

Annual emissions of CO₂ ranged from 36.3 to 40.7 ktons km⁻² y⁻¹ with a 3-year mean of 39.1 ± 2.4 ktons km⁻² y⁻¹ (Table 1). 6 7 These values are in good agreement with results from a previous measurement campaign at the BT tower in 2007 (35.5 ktons km⁻² y⁻¹-; (Helfter et al., 2011)) and London Atmospheric Emissions Inventory (LAEI) bottom-up emission estimates for the 8 9 central London boroughs of Westminster and Camden, which are the foremost spatial source areas entrained by the BT tower flux footprint. The good agreement for CO₂ obtained in the present and previous studies using different instrumentation 10 11 provides a benchmark for subsequent comparisons between top-down measurements and bottom-up inventory estimates. Due 12 to insufficient temporal coverage, individual annual budgets for 2012-2014 could not be derived for carbon monoxide. Instead, one single annual CO flux value was calculated from individual monthly averages collected in the period September 2011 – 13 14 December 2014 on the assumption that year-on-year variability was small. Furthermore, emissions of CO for August and September, when no observations were available, were estimated from a linear relation between F_{CO} and air temperature (Fig. 15 7f). The composite annual emissions estimate of 89 ± 16 t km⁻² y⁻¹ (range taken as the random uncertainty) is consistent with 16 17 the LAEI data (Table 1). Flux ratios are less sensitive to limitations in vertical transport and provide an additional means of assessing the quality of the bottom-up emission inventories and identifying poorly represented sources. Measured flux ratios 18 19 of F_{CO} to F_{CO2} were consistent with average LAEI emission ratios (Table 2, Fig. <u>\$859</u>). Measured flux ratios of F_{CO} to F_{CH4} were about half the inventoried values and measured ratios of F_{CH4} to F_{CO2} were twice the mean LAEI values (Table 2; Fig. 20 21 5758), consistent with the measured annual CH₄ fluxes (3-year mean 72 ± 3 t km⁻² y⁻¹) being more than twice the inventory 22 value. This indicates that some CH₄ sources were either underestimated or unaccounted for by the LAEI. Of the source 23 categories included in the LAEI and listed in Table 2 only gas leakage has the potential to increase the CH₄/CO₂ flux ratio, but 24 an underestimation in leakage is only a possible explanation if it follows the measured diurnal cycle, either due to changes in 25 the supply pressure or in post-meter emissions. We speculate that the diurnal, seasonal and spatial variations in F_{CH4} , and the 26 larger F_{CH4}/F_{CO2} ratio could be due a contribution of temperature-sensitive CH₄ emissions perhaps of biogenic origin (e.g. 27 increased methanogenesis from sewerage) not included in the inventories. This could explain why the net seasonal decrease in CH₄ was but half that of CO₂ (Fig. 7a and b). Previously reported discrepancies of 1.5 to > 2 between top-down and bottom-28 29 up estimates of CH₄ for the South Coast Air Basin in the greater Los Angeles (USA) area have been related to emissions from 30 landfills and other biogenic sources (Hsu et al., 2010; Wunch et al., 2009). In our study, annual methane fluxes exhibited 31 substantial spatial variability when segregated by wind sector (Fig. 8a). Fluxes of methane in the E, S and SE sectors were ca. 30% larger than the mean annual F_{CH4} estimate and exceeded the top boundary of the overall mean (taken as mean F_{CH4} + 32 maximum monthly uncertainty; Fig. 8a). In contrast, F_{CH4} from the N and NW sectors were 40% and 30% of the mean value, 33

respectively, and fell below the lower limit of the overall mean (taken as mean F_{CH4} – maximum monthly uncertainty). This 1 2 perhaps suggests more complex, spatially discrete, source distribution and composition for CH₄ compared with CO₂ and CO. 3 The linear correlation between F_{CH4} and population was strong if the highest emitting wind sectors (E, S and SE) were excluded 4 from the regression (Fig. 8b). Socio-economic temporal dynamics, such as a significant daytime influx of commuters into a 5 business district (e.g. the City of London financial district which is located 3-4 km S-SE of the BT tower), might contribute substantially to CH₄ emissions (e.g. from sewage, natural gas); in addition, the measured CH₄ emissions from such business 6 7 areas might bear no correlation with the actual resident population reported here (source: London Datastore, Greater London Authority, 2016) which can be considerably smaller than the commuting workforce. Emissions of CH_4 in the E were strongly 8 correlated with air temperature (Table 3), which suggests one or more dominant seasonal source in that wind sector. Finally, 9 10 neither test was statistically significant for emissions in the SE and S where the flux footprints entrain some of the most heavily 11 urbanised areas of central London as well as part of the river Thames. Further work is needed to investigate the potential 12 presence of additional sources of methane which might be prevalent in those wind sectors. 13 As for CH_4 , CO_2 fluxes exhibited a dependence upon air temperature in the N, NE, E and W. The seasonality of the CO_2

emissions was not statistically significant in the remaining wind sectors which might be due to the presence of substantial constant sources of CO_2 or to the prevalence of seasonal activities which do not emit CO_2 locally (e.g. more electrical heating than natural gas). However, the spatial variability of F_{CO_2} was well-captured by differences in population in the respective flux footprints of all wind sectors, except S (Fig. 8b).

18 4 Conclusions

This study presents the results of more than three years of continuous long-term eddy-covariance observations of fluxes of carbon monoxide, carbon dioxide and methane at an elevated measurement site (BT tower, 190 m a.g.l.) in central London, UK. This unique vantage point, combined with the length of the study, allowed for the spatial and temporal emission dynamics to be analysed in detail. The main conclusions are <u>that all</u> three trace gases exhibited diurnal cycles consistent with anthropogenic activities (traffic, natural gas use) <u>and</u> underwent marked seasonal dynamics, with reduced emissions in the summer.

25 <u>Seasonality of the measured fluxes</u>

Emissions of CO were strongly correlated with air temperature which is thought to be due to cold starts and reduced fuel combustion efficiency by the London fleet during the winter. Winter time emissions of CO accounted for 45% of the annual budget._Emissions of CO₂ were <u>also correlated to air temperature and were 33%</u> larger in winter than in summer.<u>and-CO₂</u> <u>emissions were predominantly controlled by the seasonal increase in natural gas consumption, although vegetation uptake</u> would also have lowered CO₂ fluxes in summer.<u>CH₄ fluxes averaged over all wind sectors decreased by 21% between winter</u> <u>and summer but unlike CO and CO₂, the correlation with air temperature was not statistically significant. averaged over all wind sectors, CH₄ fluxes decrease by 21% between winter and summer., but unlike CO and CO₂, the correlation with air</u>

- 1 temperature was not statistically significant. When segregated by wind sector, CH4 fluxes in the E and W were highstrongly
- 2 correlated with air temperature suggestive of sources with highly seasonal emission rates, possibly leaks from the natural gas
- 3 distribution network or emissions from sewage. Furthermore, CO₂ and CH₄ fluxes were positively correlated with population
- 4 density in all wind sectors except S for F_{CO2} and S, SE and E for F_{CH4}. This indicates heterogeneous source distributions and/or
- 5 densities with temporal dynamics which differ from the other wind sectors.
- 6 <u>Comparisons with atmospheric emissions inventories</u>
- 7 Measured annual emissions of CO₂ (39 ktons km⁻²) wereas in good agreement with bottom-up estimates from the London
- 8 Atmospheric Emissions Inventory (LAEI). As CO₂ is the easiest of most accurately represented of the three compounds to get
- 9 right in emission inventories, this provides confidence in the flux measurements. Similarly, the measured annual emission
- 10 budgetsbudget for CO (89 tons km⁻²) also agreed well with bottom up estimates from the London Atmospheric Emissions
- 11 Inventory (was consisting with LAEI) values which confirming confirms that the spatial distribution of the sources of this
- 12 pollutant is well captured by the inventory. <u>However, the measured</u> annual CH₄ emissions (72 tons km⁻²), however, were more
- 13 than double the LAEI value suggesting that sources are not as well-characterised by the inventory. In particular, we hypothesise
- 14 that the shortfall in inventoried CH₄ emissions can be explained by the existence of temperature-dependent sources, perhaps
- 15 of biogenic origin (e.g. sewage).
- 16

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1	References
---	------------

Andrews, G. E., Zhu, G., Li, H., Simpson, A., Wylie, J.A., Bell, M. and Tate, J. : The effect of ambient temperature on cold
start urban traffic emissions for a real world SI car, Proceedings of SAE 2004 Powertrain & Fluid Systems Conference and
Exhibition Tampa, FL, USA, October 2004, 2004.

5

Aubinet, M., Grelle, A., Ibrom, A., Rannik, U., Moncrieff, J., Foken, T., Kowalski, A. S., Martin, P. H., Berbigier, P.,
Bernhofer, C., Clement, R., Elbers, J., Granier, A., Grunwald, T., Morgenstern, K., Pilegaard, K., Rebmann, C., Snijders, W.,
Valentini, R., and Vesala, T.: Estimates of the annual net carbon and water exchange of forests: The EUROFLUX
methodology, Advances in Ecological Research, Vol 30, 30, 113-175, 2000.

10

Baldocchi, D., Falge, E., Gu, L. H., Olson, R., Hollinger, D., Running, S., Anthoni, P., Bernhofer, C., Davis, K., Evans, R.,
Fuentes, J., Goldstein, A., Katul, G., Law, B., Lee, X. H., Malhi, Y., Meyers, T., Munger, W., Oechel, W., U, K. T. P.,
Pilegaard, K., Schmid, H. P., Valentini, R., Verma, S., Vesala, T., Wilson, K., and Wofsy, S.: Fluxnet: A new tool to study the
temporal and spatial variability of ecosystem-scale carbon dioxide, water vapor, and energy flux densities, Bulletin of the
American Meteorological Society, 82, 2415-2434, 10.1175/1520-0477(2001)082<2415:fantts>2.3.co;2, 2001.

16

Baldocchi, D.: Breathing of the terrestrial biosphere: Lessons learned from a global network of carbon dioxide flux
measurement systems, Australian Journal of Botany, 56, 1-26, 10.1071/bt07151, 2008.

19

Barlow, J. F., Dunbar, T. M., Nemitz, E. G., Wood, C. R., Gallagher, M. W., Davies, F., O'Connor, E., and Harrison, R. M.:
Boundary layer dynamics over London, UK, as observed using Doppler LIDAR during REPARTEE-II, Atmospheric
Chemistry and Physics, 11, 2111-2125, 10.5194/acp-11-2111-2011, 2011.

23

Barlow, J. F., Halios, C. H., Lane, S. E., and Wood, C. R.: Observations of urban boundary layer structure during a strong
urban heat island event, Environmental Fluid Mechanics, 15, 373-398, 10.1007/s10652-014-9335-6, 2015.

26

Bjorkegren, A. B., Grimmond, C. S. B., Kotthaus, S., and Malamud, B. D.: CO₂ emission estimation in the urban environment:
Measurement of the CO₂ storage term, Atmospheric Environment, 122, 775-790,
http://dx.doi.org/10.1016/j.atmosenv.2015.10.012, 2015.

30

31 Bohnenstengel, S. I., Belcher, S. E., Aiken, A., Allan, J. D., Allen, G., Bacak, A., Bannan, T. J., Barlow, J. F., Beddows, D.

32 C. S., Bloss, W. J., Booth, A. M., Chemel, C., Coceal, O., Di Marco, C. F., Dubey, M. K., Faloon, K. H., Fleming, Z. L.,

33 Furger, M., Gietl, J. K., Graves, R. R., Green, D. C., Grimmond, C. S. B., Halios, C. H., Hamilton, J. F., Harrison, R. M., Heal,

M. R., Heard, D. E., Helfter, C., Herndon, S. C., Holmes, R. E., Hopkins, J. R., Jones, A. M., Kelly, F. J., Kotthaus, S.,
 Langford, B., Lee, J. D., Leigh, R. J., Lewis, A. C., Lidster, R. T., Lopez-Hilfiker, F. D., McQuaid, J. B., Mohr, C., Monks, P.
 S., Nemitz, E., Ng, N. L., Percival, C. J., Prevot, A. S. H., Ricketts, H. M. A., Sokhi, R., Stone, D., Thornton, J. A., Tremper,
 A. H., Valach, A. C., Visser, S., Whalley, L. K., Williams, L. R., Xu, L., Young, D. E., and Zotter, P.: Meteorology, air quality,
 and health in London the ClearfLo project, Bulletin of the American Meteorological Society, 96, 779-804, 10.1175/bams-d 12-00245.1, 2015.

8 Cambaliza M.O.L., S. P., Bogner J., Caulton D.R., Stirm B., et al.: Quantification and source apportionment of the methane
9 emission flux from the city of Indianapolis., Elementa Science of the Anthropocene, 3, 10.12952/journal.elementa.000037,
10 2015.

11

Carslaw, D. C. and Ropkins K.: openair --- an R package for air quality data analysis. Environmental Modelling & Software.
Volume 27-28, 52-61, 2012.

14

15 Carslaw D.C. and Ropkins K.: openair: Open-source tools for the analysis of air pollution data. R package version 1.716 3, http://CRAN.R-project.org/package=openair, 2016.

17

Christen, A., Coops, N. C., Crawford, B. R., Kellett, R., Liss, K. N., Olchovski, I., Tooke, T. R., van der Laan, M., and Voogt,
J. A.: Validation of modeled carbon-dioxide emissions from an urban neighborhood with direct eddy-covariance
measurements, Atmospheric Environment, 45, 6057-6069, 10.1016/j.atmosenv.2011.07.040, 2011.

21

Christen, A.: Atmospheric measurement techniques to quantify greenhouse gas emissions from cities, Urban Climate, 10, Part
 2, 241-260, http://dx.doi.org/10.1016/j.uclim.2014.04.006, 2014.

24

Crosson, E. R.: A cavity ring-down analyzer for measuring atmospheric levels of methane, carbon dioxide, and water vapor,
Applied Physics B-Lasers and Optics, 92, 403-408, 10.1007/s00340-008-3135-y, 2008.

27

Evans, S.: 3D cities and numerical weather prediction models: An overview of the methods used in the LUCID project,
available at http://discovery.Ucl.Ac.Uk/17404/1/17404.Pdf UCL Working Paper Series, 2009.

30

Fiddler, M. N., Begashaw, I., Mickens, M. A., Collingwood, M. S., Assefa, Z., and Bililign, S.: Laser spectroscopy for
atmospheric and environmental sensing, Sensors, 9, 10447-10512, 10.3390/s91210447, 2009.

- 1 Finkelstein, P. L., and Sims, P. F.: Sampling error in eddy correlation flux measurements, Journal of Geophysical Research:
- 2 Atmospheres, 106, 3503-3509, 10.1029/2000jd900731, 2001.
- 3
- Foken, T., and Wichura, B.: Tools for quality assessment of surface-based flux measurements, Agricultural and Forest
 Meteorology, 78, 83-105, 10.1016/0168-1923(95)02248-1, 1996.
- 6
- 7 Foken, T.: Micrometeorology, Springer-Verlag Berlin Heidelberg, 308 pp., 2008.
- Foken, T., Gödecke, M., Mauder, M., Mahrt, L., Amiro, B., and Munger, W.: Post-field data quality control, in: Handbook of
 micrometeorology, edited by: Lee, X., Kluwer Academic Publishers, 2004.
- 10
- Gioli, B., Toscano, P., Lugato, E., Matese, A., Miglietta, F., Zaldei, A., and Vaccari, F. P.: Methane and carbon dioxide fluxes
 and source partitioning in urban areas: The case study of Florence, Italy, Environmental Pollution, 164, 125-131,
 10.1016/j.envpol.2012.01.019, 2012.
- 14
- 15 Greater London Authority, London Datastore: http://data.london.gov.uk/, 2016.
- 16
- Grimmond, C. S. B., and Christen, A.: Flux measurements in urban ecosystems, in: FluxLetter, The newsletter of FLUXNET,
 1, FLUXNET, 2012.
- 19
- Halios, C. H., and Barlow, J. F.: Observations of the morning development of the urban boundary layer over London, UK,
 taken during the actual project., Boundary-Layer Meteorology, under review 2016.
- 22
- Harrison, R. M., Dall'Osto, M., Beddows, D. C. S., Thorpe, A. J., Bloss, W. J., Allan, J. D., Coe, H., Dorsey, J. R., Gallagher,
 M., Martin, C., Whitehead, J., Williams, P. I., Jones, R. L., Langridge, J. M., Benton, A. K., Ball, S. M., Langford, B., Hewitt,
 C. N., Davison, B., Martin, D., Petersson, K. F., Henshaw, S. J., White, I. R., Shallcross, D. E., Barlow, J. F., Dunbar, T.,
 Davies, F., Nemitz, E., Phillips, G. J., Helfter, C., Di Marco, C. F., and Smith, S.: Atmospheric chemistry and physics in the
 atmosphere of a developed megacity (London): an overview of the REPARTEE experiment and its conclusions, Atmospheric
 Chemistry and Physics, 12, 3065-3114, 2012.
- 29
- Helfter, C., Famulari, D., Phillips, G. J., Barlow, J. F., Wood, C. R., Grimmond, C. S. B., and Nemitz, E.: Controls of carbon
 dioxide concentrations and fluxes above central London, Atmospheric Chemistry and Physics, 11, 1913-1928, 10.5194/acp11-1913-2011, 2011.
- 33

Hsu, Y.-K., VanCuren, T., Park, S., Jakober, C., Herner, J., FitzGibbon, M., Blake, D. R., and Parrish, D. D.: Methane
 emissions inventory verification in Southern California, Atmospheric Environment, 44, 1-7, 10.1016/j.atmosenv.2009.10.002,
 2010.

5 International Energy Agency: World energy outlook, http://www.iea.org/publications/freepublications/publication/world6 energy-outlook-2012.html, 2012.

7

8 IPCC (International Panel on Climate Change): IPCC fifth assessment report: Climate change 2013, 2013.

9

Jarvi, L., Nordbo, A., Junninen, H., Riikonen, A., Moilanen, J., Nikinmaa, E., and Vesala, T.: Seasonal and annual variation
of carbon dioxide surface fluxes in Helsinki, Finland, in 2006-2010, Atmospheric Chemistry and Physics, 12, 8475-8489,
10.5194/acp-12-8475-2012, 2012.

13

Kormann, R., and Meixner, F. X.: An analytical footprint model for non-neutral stratification, Boundary-Layer Meteorology,
99, 207-224, 10.1023/a:1018991015119, 2001.

16

17 Kotthaus, S., and Grimmond, C. S. B.: Identification of micro-scale anthropogenic CO₂, heat and moisture sources - processing 18 eddy covariance fluxes for а dense urban environment, Atmospheric Environment, 57. 301-316, 19 10.1016/j.atmosenv.2012.04.024, 2012.

20

Kotthaus, S., and Grimmond, C. S. B.: Energy exchange in a dense urban environment – part II: Impact of spatial heterogeneity
of the surface, Urban Climate, 10, Part 2, 281-307, http://dx.doi.org/10.1016/j.uclim.2013.10.001, 2014a.

23

Kotthaus, S., and Grimmond, C. S. B.: Energy exchange in a dense urban environment – part I: Temporal variability of longterm observations in central London, Urban Climate, 10, Part 2, 261-280, http://dx.doi.org/10.1016/j.uclim.2013.10.002,
2014b.

27

Langford, B., Nemitz, E., House, E., Phillips, G. J., Famulari, D., Davison, B., Hopkins, J. R., Lewis, A. C., and Hewitt, C.
N.: Fluxes and concentrations of volatile organic compounds above central London, UK, Atmospheric Chemistry and Physics,
10, 627-645, 10.5194/acp-10-627-2010, 2010.

31

Liu, H. Z., Feng, J. W., Jarvi, L., and Vesala, T.: Four-year (2006-2009) eddy covariance measurements of CO₂ flux over an
urban area in Beijing, Atmospheric Chemistry and Physics, 12, 7881-7892, 10.5194/acp-12-7881-2012, 2012.

Lowry, D., Holmes, C. W., Rata, N. D., O'Brien, P., and Nisbet, E. G.: London methane emissions: Use of diurnal changes in
 concentration and delta C-13 to identify urban sources and verify inventories, Journal of Geophysical Research-Atmospheres,
 106, 7427-7448, 10.1029/2000jd900601, 2001.

4

McKain, K. K., Down, A., Raciti, S. M., Budney, J., Hutyra, L. R., Floerchinger, C., Herndon, S. C., Nehrkorn, T., Zahniser,
M. S., Jackson, R. B., Phillips, N., and Wofsy, S. C.: Methane emissions from natural gas infrastructure and use in the urban
region of Boston, Massachusetts, Proceedings of the National Academy of Sciences of the United States of America, 112,
1941-1946, 10.1073/pnas.1416261112, 2015.

9

Nam, E. K., Jensen, T. E., and Wallington, T. J.: Methane emissions from vehicles, Environmental Science & Technology, 38,
2005-2010, 10.1021/es034837g, 2004.

12

Moncrieff, J., Clement, R., Finnigan, J., and Meyers, T.: Averaging, detrending and filtering of eddy covariance time series.
In: Handbook of Micrometeorology, Lee, X. (Ed.), Kluwer Academic Publishers, 2004.

15

16 O'Shea, S. J., Allen, G., Fleming, Z. L., Bauguitte, S. J. B., Percival, C. J., Gallagher, M. W., Lee, J., Helfter, C., and Nemitz,

17 E.: Area fluxes of carbon dioxide, methane, and carbon monoxide derived from airborne measurements around greater London:

18 A case study during summer 2012, Journal of Geophysical Research-Atmospheres, 119, 4940-4952, 10.1002/2013jd021269,
19 2014.

20

Mayor of London Office: London population confirmed at record high: https://www.london.gov.uk/media/mayor-press releases/2015/02/london-population-confirmed-at-record-high, 2015.

23

Pawlak, W., Fortuniak, K., and Siedlecki, M.: Carbon dioxide flux in the centre of Lodz, Poland - analysis of a 2-year eddy
covariance measurement data set, International Journal of Climatology, 31, 232-243, 10.1002/joc.2247, 2011.

26

Pawlak, W. and Fortuniak, K.: Eddy covariance measurements of the net turbulent methane flux in the city centre – results of
2 years campaign in Łódź, Poland, Atmos. Chem. Phys. Discuss., 2016, 1-38, 2016.

29

Peltola, O., Hensen, A., Helfter, C., Marchesini, L. B., Bosveld, F. C., van den Bulk, W. C. M., Elbers, J. A., Haapanala, S.,
Holst, J., Laurila, T., Lindroth, A., Nemitz, E., Rockmann, T., Vermeulen, A. T., and Mammarella, I.: Evaluating the
performance of commonly used gas analysers for methane eddy covariance flux measurements: The InGOS inter-comparison
field experiment, Biogeosciences, 11, 3163-3186, 10.5194/bg-11-3163-2014, 2014.

1	United Nations:	World urbanization	ation prospect	s. http://esa.u	n.org/unpd/w	up/highlights/wu	p2014-highlights.p	df. 2014.

3	Velasco, E., and Roth, M.: Cities as net sources of co2: Review of atmospheric CO2 exchange in urban environments measured
4	by eddy covariance technique. Geography Compass 4, 1238-1259, 10, 1111/j, 1749-8198, 2010, 00384 x, 2010.

2

Ward, H. C., Kotthaus, S., Grimmond, C. S. B., Bjorkegren, A., Wilkinson, M., Morrison, W. T. J., Evans, J. G., Morison, J.
I. L., and Iamarino, M.: Effects of urban density on carbon dioxide exchanges: Observations of dense urban, suburban and
woodland areas of southern England, Environmental Pollution, 198, 186-200, 10.1016/j.envpol.2014.12.031, 2015.

9

Wennberg, P. O., Mui, W., Wunch, D., Kort, E. A., Blake, D. R., Atlas, E. L., Santoni, G. W., Wofsy, S. C., Diskin, G. S.,
Jeong, S., and Fischer, M. L.: On the sources of methane to the Los Angeles atmosphere, Environmental Science &
Technology, 46, 9282-9289, 10.1021/es301138y, 2012.

13

14 Wood, C. R., Lacser, A., Barlow, J. F., Padhra, A., Belcher, S. E., Nemitz, E., Helfter, C., Famulari, D., and Grimmond, C. S.

B.: Turbulent flow at 190 m height above London during 2006-2008: A climatology and the applicability of similarity theory,
Boundary-Layer Meteorology, 137, 77-96, 10.1007/s10546-010-9516-x, 2010.

17

Wunch, D., Wennberg, P. O., Toon, G. C., Keppel-Aleks, G., and Yavin, Y. G.: Emissions of greenhouse gases from a North
American megacity, Geophysical Research Letters, 36, 10.1029/2009gl039825, 2009.

20

Zazzeri, G., Lowry, D., Fisher, R. E., France, J. L., Lanoiselle, M., and Nisbet, E. G.: Plume mapping and isotopic
characterisation of anthropogenic methane sources, Atmospheric Environment, 110, 151-162,
10.1016/j.atmosenv.2015.03.029, 2015.

1 Tables

2 Table 1: Annual totals of carbon dioxide flux and methane flux calculated from monthly averages for the period 2012-2014. The 3 data period September 2012-March 2013 (no ultrasonic anemometer) was gapfilled using available monthly averages obtained over 4 the remaining measurement period. Due to insufficient temporal coverage, individual annual budgets for 2012-2014 could not be 5 derived for the carbon monoxide flux. A composite annual emissions estimate was compiled instead which makes use of all available 6 monthly averages of F_{CO} over the study period September 2011 to December 2014. Data from the London Atmospheric Emissions 7 Inventory (LAEI; emissions for the central London boroughs of Westminster and Camden) and previous measurement campaigns 8 are provided for comparison with the current study.

9

	Reference	FCO ₂	FCH ₄	FCO
		[kt km ⁻²]	[t km ⁻²]	[t km ⁻²]
2012	This study	40.2	69	-
2013	This study	40.7	75	-
2014	This study	36.3	72	-
Mean ± SD	This study	39.1 ± 2.4	72 ± 3	89
Random	This study	6.5	12	16
uncertainty				
Emissions	LAEI	38.7	29	110
inventory				
(2012)				
London 2007	(Helfter et al., 2011)	35.5	-	-
London	(Ward et al., 2015)	46.6		
London	(Harrison et al., 2012)			150 - 220
Autumn				
2007/08				
London July	(O'Shea et al., 2014) ⁺	29.0	66	106
2012				

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12

- 1 Table 2: Emission ratios from measurements and the London Atmospheric Emissions Inventory (LAEI). Measured quantities are
- 2 mean, median and range of monthly emissions segregated by wind direction.

Emission category	Zone	FCH4/FCO2	Fco/Fco2	F CO/ F CH4
Measured (this study):				
Mean		0.0019	0.0018	0.9739
Median		0.0019	0.0021	1.1304
Minimum		0.0017	0.0004	0.1972
Maximum		0.0022	0.0027	1.5951
LAEI (all sources)	Central	0.0009	0.0039	4.1023
	Inner	0.0010	0.0024	2.4546
	Outer	0.0094	0.0018	0.1965
Domestic Coal	Central	-	-	-
	Inner	0.0020	0.0460	22.400
	Outer	0.0020	0.0460	22.414
Domestic Oil	Everywhere	0.0001	0.0006	4.0000
Domestic Gas	Everywhere	0.0001	0.0006	6.0375
Non-Domestic Gas	Everywhere	0.0001	0.0002	2.2642
Boilers	Central	0.0001	4E-05	0.3270
	Inner	0.0001	5E-05	0.3548
	Outer	0.0001	5E-05	0.3482
Gas Leakage	Everywhere	26.607	-	-
Non-Road Mobile	Central	0.0002	0.0347	213.55
Machinery, agriculture & other				
	Inner	0.0003	0.0377	119.75
	Outer	0.3525	0.0576	0.1633
Road - All Sources	Central	-	0.0021	-
	Inner	-	0.0013	-
	Outer	-	0.0013	-

1 Table 3: Goodness of fit of the linear regression between wind sector-segregated monthly methane fluxes (F_{CH4}), carbon dioxide

fluxes (F_{CO2}) and monthly mean air temperature (T_{air}). The superscripts (-) and (+) denote the sign of slope for each linear regression.

P-values in **bold** denote statistical significance.

R ²	Ν	NE	Е	SE	S	SW	W	NW ⁴
F _{CH4} v. T _{air}	0.55(-)	0.43(-)	0.73(-)	0.08(-)	0.04(-)	0.05(-)	0.60(-)	0.21(-)
p-value	0.0060	0.0197	0.0004	0.3627	0.5150	0.4790	0.0033	0.1367 5
FCO2 V. Tair	0.51 ⁽⁻⁾	0.69(-)	0.80(-)	0.19(-)	0.18(-)	0.27(-)	0.60 ⁽⁻⁾	0.34(-) 6
p-value	0.0203	0.0021	0.0003	0.1496	0.1984	0.1650	0.0081	0.0776 7
								8

1 Figures







5 Figure 1: Frequency of occurrence of x₉₀ (distance from the tower where 90% of the measured fluxes originated from) centred at 6 the BT tower as a function of wind direction and season. The flux footprint was estimated using an analytical model for nonear-7 neutral stratification (Kormann and Meixner, 2001) and the plots were produced using the openair package for R (Carslaw and 8 Ropkins, 2012; Carslaw and Ropkins, 2016). Bin dimensions: 10° (angular scale) × 1000 m (radial scale). Data period 15/09/2011-9 31/12/2014.





Figure 2: Normalised cospectra of T (sonic temperature), CO₂ and CH₄ with respect to w (vertical wind component). Each
cospectrum is an average of 24 half-hourly cospectra (data period 12/03/2013 7:00 - 18:00). Regression of spectra for frequencies >
0.1 Hz marked by solid line for Co(wT) and dashed line for both Co(wCO₂) and Co(wCH₄).



Figure 3: Comparison of half-hourly fluxes of CO₂ measured in March, August and October 2013 by a closed-path Picarro G1301f operating at 1 Hz following high-frequency loss correction and an open-path Li7500 analyser operating at 20 Hz at the top of BT
tower (sensor height: 190 m a.g.l.). Dashed line is 1:1 line.



Figure 4: Mean diurnal profiles of CO₂ fluxes for (a) winter (DJF), and (b) summer (JJA) for the data period 15/09/2011 – 31/12/2013;
(c) CH4 fluxes in summer observed at the BT tower site (190 m a.g.l.) and the KCL site (50 m a.g.l.) over the period 19/08 – 01/10/2015;
mixing height obtained from Doppler LIDAR measurements for (d) winter and (e) summer (Bohnenstengel et al., 2015); road traffic
counts (f) winter and (g) summer (average of 246 counting stations distributed throughout the London conurbation; source:
Transport for London, 2012 data). Fluxes are normalised (subtraction of diurnal minimum and normalisation by offset-corrected maximum value) to aid visualisation and comparison of the temporal dynamics
The shaded areas represent the 95% confidence interval.



Figure 5: Comparison between <u>average</u> diurnal <u>average-profiles of</u> CO₂ fluxes measured at the BT tower (190 m a.g.l.) and at the
 KCL site (50 m a.g.l.) in the period 15/09/2011 – 31/12/2013, separated into eight wind-direction sectors based on the wind direction
 observed at BT tower.





Figure 6: Fluxes of (a, d, g) methane (F_{CH4}), (b, e, h) carbon dioxide (F_{CO2}) and (c, f, i) carbon monoxide (F_{CO}) observed at BT tower
with a closed-path gas analyser (from 15/09/2011 to 31/12/2014): (a)-(c) mean diurnal patterns with 95% confidence interval
(shading), (d)-(f) as (a)-(c) but segregated into wind sectors and (g)-(i) by day of the week.



Figure 7: (a)-(c): Monthly averages of F_{CH4}, F_{CO2} and F_{CO} (September 2011-December 2014); (d)-(f): F_{CH4}, F_{CO2} and F_{CO} as a function
of monthly mean air temperature. Solid lines are linear regressions and shaded areas are 95% confidence intervals. No F_{CO}
measurements were available in August and September due to instrument downtime.



2

3 Figure 8: Annual fluxes of carbon dioxide (F_{C02}) and methane (F_{CH4}) measured by eddy-covariance at the BT tower in central 4 London as a function of (a) wind direction; solid lines are mean annual emissions (2012-2014) without wind sector segregation. Mean 5 measurement uncertainty (taken as the maximum of monthly uncertainties for each gas) is denoted by a blue (F_{CO2}) and red striped 6 areas (F_{CH4}). (b) Data from plot (a) as a function of population within each wind sector-specific flux footprint area. The spatial extent 7 of the footprint for each wind sector was derived from footprint statistics (Fig. 1) with the approximation that the typical extent was 8 of the order of 10 km for NE-SW and 15 km for W-N. Population data (source: London Datastore, Greater London Authority, 2016) 9 are on a ward basis (i.e. sub-borough administrative unit). Linear regression (dashed lines), with exclusion of S sector data point for 10 FCO2, E, S and SE for FCH4 (identified by their wind sector abbreviations). NB: FCO2 and associated uncertainty are divided by 1000 11 to aid visualisation.