

1 **Response to Referee # 3**

2

3 We thank the reviewer for providing provocative and helpful comments.

4

5 **Reviewer Comments**

6

7 *The paper has improved through the review process, but I still have a significant concern that*
8 *the authors draw conclusions to eliminate the possibility of a large underestimated NG source*
9 *that are not fully supported by the evidence in the paper. On the topic of whether or not there is*
10 *evidence for a significant missing NG source in Indianapolis, the authors seem to be posing the*
11 *wrong the question. The relevant question is not whether NG or the SSLF is the largest source in*
12 *Indianapolis. No one disputes that there is a large landfill source in the middle of Indianapolis.*
13 *The question which is relevant to the existing literature and society is whether NG emissions are*
14 *significantly underestimated by the inventory. Can you reject the hypothesis that the 40%*
15 *difference between top-down and bottom-up that remains after the domain issue is fixed is due*
16 *to NG leaks? This question needs to be clearly and consistently addressed in the text. At a*
17 *minimum, I would like to see the following statements in the paper revised to accurately and*
18 *consistently represent the evidence given.*

19

20 We do not have a fundamental disagreement with the reviewer on these points.

21

22 We had some errors in our percentage calculations, which are now fixed.

23

24 We cannot “reject the hypothesis that the 40% (it is 91% actually) difference between top-
25 down and bottom-up that remains after the domain issue is fixed is due to NG leaks.” We did
26 not intend to make this point, and we have clarified the text to make this clear.

27

28 We do conclude that the discrepancy between top-down and bottom-up is less than that
29 suggested by the Lamb et al., (2016) paper. We also do show evidence for the lack of large,
30 persistent point sources other than the SSLF. We have revised the text to clarify these
31 conclusions, and to make sure that these are not conflated.

32

33 We also conclude that, given the large potential for random error in the background (one main
34 focus of this manuscript), we cannot yet be confident in the significance of the difference
35 between the inventory and the atmospheric estimate of the diffuse NG source. We have gone
36 through the text to make this point clear where appropriate.

37

38 *Abstract, Line 43: “Leaks from the natural gas distribution system that were detected with the*
39 *tower network appeared localized and non-permanent and do not appear to constitute as large*
40 *of a source of CH4 as previously hypothesized by some top-down studies.” It sounds like you are*
41 *providing evidence to reject the findings of previous top-down studies, but you are actually only*
42 *providing evidence for the absence of point sources rather than a diffuse source comprised of*
43 *many small point sources. This sentence needs rephrasing to more fairly represent the evidence*
44 *given in the paper.*

45
46 Now we say:
47
48 The data from the towers confirm that the strongest CH₄ source in Indianapolis is South Side
49 Landfill. Leaks from the natural gas distribution system that were detected with the tower
50 network appeared localized and non-permanent. Our simple atmospheric budget analyses
51 estimate magnitude of the diffuse NG source to be 70% higher than inventory estimates, but
52 more comprehensive analyses are needed.

53
54
55 *Line 463: Please give a number for "somewhat"*

56
57 Now we say:
58
59 "If we assume that SSLF emissions are generally known (GHG reporting program) that would
60 indicate that emissions from NG distribution are likely to be about 14 mol/s (70%) higher than
61 what both of the inventories currently estimate but within the error bars of Lamb et al., (2016)'s
62 inventory calculation."

63
64 *Line 500: "Thus, the diffuse NG source suspected to be twice as large as the SSLF source (Lamb*
65 *et al., 2016) does not appear to be supported by these data." This is true for large point sources*
66 *but not for a broad diffuse source. Please reword or add a sentence to reflect such a possibility.*

67
68 Here is the reworked part of that paragraph:

69
70 "None of the individual leaks appears to be similar in magnitude to the emissions that originate
71 from SSLF. Diffuse NG emissions comparable to the SSLF source (Lamb et al., 2016) may
72 exist. Our flux estimations at towers 8 and 13, however, imply that the magnitude of NG diffuse
73 source suggested by the top-down analyses in Cambaliza et al. (2015) and Lamb et al. (2016) are
74 probably overestimates (see section 3.3). We hypothesize that the relatively high Indianapolis
75 CH₄ emissions (see Fig. 1) reported by Cambaliza et al. (2015) could be a result of random errors
76 in upwind conditions (see section 3.2) influencing the small number of airborne flux estimates."

77
78 *Line 574: "Analysis of the INFLUX observation data suggests that inventories for Indianapolis are*
79 *mostly accurate and that there is no clear evidence of a large, diffuse NG source of CH₄ as*
80 *implied by Lamb et al. (2016)." I take issue with the "mostly accurate" phrasing since you still*
81 *have a significant difference between top-down and bottom-up (Line 369).*

82
83 Changed to:

84
85 "Analysis of the INFLUX tower observations suggest a diffuse NG source that exceeds both of
86 the inventory estimates by 70%, but additionally our analysis shows that the discrepancy is less
87 than that proposed by highest values reported in Lamb et al. (2016) (see Fig. 1). Uncertainty
88 remains regarding the magnitude of the diffuse NG source of CH₄. The only major point source

89 in the city is SSLF and it is observed at multiple towers. There is an evidence for occasional
90 point-source NG leaks, but they appear to be transient in time, and limited in their strength.”

91

92 *Other comments:*

93

94 *Abstract, Line 47: I do not understand the meaning of “real”. Please rephrase.*

95

96 Now it says:

97

98 “Long-term averaging, spatially-extensive upwind mole fraction observations, mesoscale
99 atmospheric modeling of the regional emissions environment, and careful treatment of the times
100 of day are recommended for precise and accurate quantification of urban CH₄ emissions.”

101

102 *Line 27: Suggest rewording “...and (4) the presence of unknown CH₄ sources.” -> “...and (4) CH₄*
103 *sources that are not accounted for in the inventory” or “...and (4) CH₄ source types that are*
104 *absent from the inventory”*

105

106 Done.

107

108 *Line 30: Suggest rewording “...about 35% and thereby lessens the discrepancy by bringing total*
109 *city flux within the error range of one of the two inventories.” -> “...about 35%, thereby*
110 *lessening the discrepancy and bringing total city flux within the error range of one of the two*
111 *inventories.”*

112

113 Done.

114

115 *Line 125: Not clear what “both” refers to*

116

117 Changed to:

118

119 “...temporal variability in urban emissions, which is not captured by the existing top-down
120 studies...”

121

122 *Line 139-140: suggest: “and boundary layer depth compared to nearby rural areas...”*

123

124 Done.

125

126 *Line 494: Why do you think the plumes are from the residential sector? State your evidence or*
127 *else remove.*

128

129 Changed to, “Because no other tower sees these enhancements (at least at comparable
130 magnitudes), we believe that they are the result of nearby NG leaks.”

131

132 **Background Heterogeneity and Other Uncertainties in**
133 **Estimating Urban Methane Flux: Results from the**
134 **Indianapolis Flux (INFLUX) Experiment**

135
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150 Davis (kjd10@psu.edu)

151 **Abstract**

152 As natural gas extraction and use continues to increase, the need to quantify emissions of
153 methane (CH₄), a powerful greenhouse gas, has grown. Large discrepancies in Indianapolis CH₄
154 emissions have been observed when comparing inventory, aircraft mass-balance, and tower
155 inverse modeling estimates. Four years of continuous CH₄ mole fraction observations from a
156 network of nine towers as a part of the Indianapolis Flux Experiment (INFLUX) are utilized to
157 investigate four possible reasons for the abovementioned inconsistencies: (1) differences in
158 definition of the city domain, (2) a highly temporally variable and spatially non-uniform CH₄
159 background, (3) temporal variability in CH₄ emissions, and (4) CH₄ sources that are not
160 accounted for in the inventory. Reducing the Indianapolis urban domain size to be consistent
161 with the inventory domain size decreases the CH₄ emission estimation of the inverse modeling
162 methodology by about 35%, thereby lessening the discrepancy and bringing total city flux within
163 the error range of one of the two inventories. Nevertheless, the inverse modeling estimate still

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170 | remains about 91% higher than inventory estimates. Hourly urban background CH₄ mole
171 | fractions are shown to be spatially heterogeneous and temporally variable. Variability in
172 | background mole fractions observed at any given moment and a single location could be up to
173 | about 50 ppb depending on a wind direction, but decreases substantially when averaged over
174 | multiple days. Statistically significant, long-term biases in background mole fractions of 2-5 ppb
175 | are found from single point observations for most wind directions. Boundary layer budget
176 | estimates suggest that Indianapolis CH₄ emissions did not change significantly when comparing
177 | 2014 to 2016. However, it appears that CH₄ emissions may follow a diurnal cycle with daytime
178 | emissions (12-16 LST) approximately twice as large as nighttime emissions (20-5 LST). We
179 | found no evidence for large CH₄ point sources that are otherwise missing from the inventories.
180 | The data from the towers confirm that the strongest CH₄ source in Indianapolis is South Side
181 | Landfill. Leaks from the natural gas distribution system that were detected with the tower
182 | network appeared localized and non-permanent. Our simple atmospheric budget analyses
183 | estimate magnitude of the diffuse NG source to be 70% higher than inventory estimates, but
184 | more comprehensive analyses are needed. Long-term averaging, spatially-extensive upwind
185 | mole fraction observations, mesoscale atmospheric modeling of the regional emissions
186 | environment, and careful treatment of the times of day are recommended for precise and accurate
187 | quantification of urban CH₄ emissions.

188

189 | 1 Introduction

190 | From the beginning of the Industrial Revolution to 2011, atmospheric methane (CH₄) mole
191 | fractions increased by a factor of 2.5 due to anthropogenic processes such as fossil fuel
192 | production, waste management, and agricultural activities (Ciais et al., 2013). The increase in

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209 CH₄ is a concern as it is a potent greenhouse gas (GHG) with a global warming potential 28-34
210 times greater than that of CO₂ over a period of 100 years (Myhre et al., 2013). The magnitudes
211 of component CH₄ sources, and the causes of variability in the global CH₄ budget are not well
212 understood although there is some evidence that biogenic emissions may play an important role
213 in the recent CH₄ increases (Nisbet et al., 2016; Saunio et al., 2016). Improved understanding
214 of CH₄ emissions is needed (National Academies of Sciences and Medicine, 2018).

215 In particular, the estimates of continental U.S. anthropogenic CH₄ emissions disagree.
216 Inventories from Environment Protection Agency (EPA) and Emissions Database for Global
217 Atmospheric Research (EDGAR) in 2008 reported emission values of 19.6 and 22.1 TgC y⁻¹
218 (U.S. EPA, 2013; European Commission Joint Research Centre and Netherlands Environmental
219 Assessment Agency, 2010). However, top-down methodologies using aircraft and inverse
220 modeling framework found emission values of 32.4 ± 4.5 TgC y⁻¹ for 2004 and 33.4 ± 1.4 TgC
221 y⁻¹ for 2007-2008 respectively (Kort et al., 2008; Miller et al., 2013). Underestimation of natural
222 gas (NG) production and agricultural sources are possible reasons for this disagreement (Miller
223 et al., 2013; Brandt et al., 2014; Jeong et al., 2014). Efforts to reconcile GHGs emissions
224 estimates using atmospheric methods and inventory assessment have sometimes succeeded
225 (Schuh et al., 2013; Zavala-Araiza et al., 2015; Turnbull et al., 2019) when careful attention is
226 given to the details of each method, and targeted atmospheric data are available. A recent
227 synthesis of emissions from the U.S. NG supply chain demonstrated similar success and
228 concluded that current inventory estimates of emissions from U.S. NG production are too low
229 and that emission from NG distribution is one of the greatest remaining sources of uncertainty in
230 the NG supply chain (Alvarez et al., 2018).

231 Due to the uncertainties in CH₄ emissions from NG distribution it is natural that urban
232 emissions are of interest as well. For example, two studies (McKain et al., 2015; Hendrick et al.,
233 2016) indicate that ~60-100% of Boston CH₄ emissions are attributable to the NG distribution
234 system. Recent studies of urban CH₄ emissions in California indicate that the California Air
235 Resources Board (CARB) inventory tends to underestimate the actual CH₄ urban fluxes possibly
236 due to fugitive emissions from NG infrastructures in urban environments (Wunch et al., 2009;
237 Jeong et al., 2016; Jeong et al., 2017). The accuracy and precision of atmospheric estimates of
238 urban CH₄ emissions are limited by available atmospheric observations (Townsend-Small et al.,
239 2012), potential source magnitude variability with time (Jackson et al., 2014; Lamb et al., 2016),
240 errors in atmospheric transport modeling (Hendrick et al., 2016; Deng et al., 2017; Sarmiento et
241 al., 2017), and complexity in atmospheric background conditions (Cambaliza et al., 2014; Karion
242 et al., 2015; Heimbürger et al., 2017). In this work, detailed analysis of urban CH₄ mole
243 fractions is performed for the city of Indianapolis to better understand the aforementioned
244 uncertainties of urban CH₄ emissions.

245 The Indianapolis Flux Experiment (INFLUX; Davis et al., 2017) is a testbed for
246 improving quantification of urban GHGs emissions and their variability in space and time.
247 INFLUX (<http://influx.psu.edu>) is located in Indianapolis partly because of its isolation from
248 other urban centers and the flat Midwestern terrain. It includes a very dense GHGs monitoring
249 network, comprised of irregular insitu aircraft measurements (Heimbürger et al., 2017;
250 Cambaliza et al., 2014), continuous in situ observations from communications towers using
251 cavity ring-down spectroscopy (Richardson et al., 2017; Miles et al., 2017), and automated flask
252 sampling systems for quantification of a wide variety of trace gases (Turnbull et al., 2015).
253 Meteorological sensors include a Doppler lidar providing continuous boundary layer depth and

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255 wind profiles, and tower-based eddy covariance measurements of the fluxes of momentum,
256 sensible and latent heat (Sarmiento et al., 2017). The network is designed for emissions
257 quantification using top-down methods such as tower-based inverse modeling (Lauvaux et al.,
258 2016) and aircraft mass balance estimates (Cambaliza et al., 2015).

259 Lamb et al. (2016) compared Indianapolis CH₄ emissions estimates from a variety of
260 approaches, specifically inventory, aircraft mass balances, and inverse modeling. The results
261 revealed large mean differences among the city fluxes estimated from these methods (Fig. 1). In
262 general, the inventory methods arrived at lower estimates of emissions compared to the
263 atmospheric, or top-down approaches. CH₄ fluxes calculated using the aircraft mass balance
264 technique varied considerably between flights, more than would be expected from propagation of
265 errors of the component measurements (Cambaliza et al., 2014; Lamb et al., 2016). The
266 atmospheric inverse estimate was significantly higher than the inventory and some of the
267 aircraft-derived values.

268 Biogenic emissions from the city are dominated by a landfill close to downtown, and
269 these emissions are thought to be fairly well known (GHG reporting program). Although
270 evidence of possible variability in landfill emissions exists from Cambaliza et al. (2015), which
271 used aircraft mass balance on five different occasions to calculate CH₄ flux from this landfill.
272 Uncertainty in total city emissions is mainly driven by the uncertainty in thermogenic emissions,
273 which are hypothesized to emerge largely from the NG distribution system (Mays et al., 2009;
274 Cambaliza et al., 2015; Lamb et al., 2016). In this study, we explore potential explanations for
275 the discrepancies in CH₄ emissions estimates from Indianapolis and posit methods and
276 recommendations for the study of CH₄ emissions from other urban centers.

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278 We examine four different potential explanations for the CH₄ flux discrepancies reported
279 in Lamb et al. (2016): (1) inconsistent geographic boundaries between top-down and bottom-up
280 studies, (2) heterogeneity in the urban scale CH₄ background and (3) temporal variability in
281 urban emissions, ~~which is not captured by the existing~~ top-down studies, and (4) CH₄ sources
282 that are not accounted for in the inventories. Well-calibrated CH₄ sensors on the INFLUX tower
283 network (Miles et al., 2017) collected continuous CH₄ observations from 2013 to 2016 and
284 provide a unique opportunity to explore these issues.

285

286 2 Methods

287

288 2.1 Experimental site

289 This study uses data from a tower-based GHG observational network located in the city and
290 surrounding suburbs of Indianapolis, Indiana in the Midwestern U.S. Prior studies have used
291 varying definitions for the region of Indianapolis (Cambaliza et al., 2015, Lamb et al., 2016). In
292 this work, we follow Gurney et al. (2012) and define Indianapolis as the area of Marion County.
293 The flat terrain of the region simplifies interpretation of the atmospheric transport. The land-
294 surface heterogeneity inherent in the urban environment (building roughness, spatial variations in
295 the surface energy balance) does have a modest influence on the ~~wind and boundary layer depth~~
296 ~~within the city compared to nearby rural areas~~ (Sarmiento et al., 2017).

297 Figure 2 shows two domains that have been used for the evaluation of Indianapolis CH₄
298 emissions (Lamb et al., 2016; Lauvaux et al., 2016). The first domain is the whole area shown in
299 the figure enclosing both Indianapolis and places that lie outside of its boundaries. This domain
300 was used for the inversion performed in Lamb et al. (2016). The second domain is Marion

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309 County outlined with a green dashed line. It is assumed here that this domain is much more
310 representative of the actual Indianapolis municipal boundaries as this area encompasses the
311 majority of the urban development associated with the city of Indianapolis (Gurney et al., 2012).
312 The larger domain has three additional landfills that based on the EPA gridded inventory
313 (Maasackers et al., 2016) increase Indianapolis CH₄ emissions by about 50% when compared to
314 the smaller domain. The inversion explained in Lamb et al. (2016) has been rerun for two of the
315 domains mentioned above and the results (Fig. 1) have been reexamined.

316

317 **2.2 INFLUX tower network**

318 The continuous GHG measurements from INFLUX are described in detail in Richardson et al.
319 (2017). The measurements were made using wavelength-scanned cavity ring down
320 spectrometers (CRDS, Picarro, Inc., models G2301, G2302, G2401, and G1301), installed at the
321 base of existing communications towers, with sampling tubes secured as high as possible on each
322 tower (39 – 136 m above ground level (AGL); Miles et al., 2017). A few towers also included
323 measurements at 10 m AGL and one or two intermediate levels. While INFLUX tower in-situ
324 measurements began in September 2010, here we focus on the CH₄ measurements from 2013 –
325 2016. From June through December 2012, there were two or three towers with operational CH₄
326 measurements. By July 2013, five towers included measurements of CH₄, and throughout the
327 majority of the years 2015 – 2016 there were eight INFLUX towers with CH₄ measurements
328 (Fig. 3). Flask to in-situ comparisons and round-robin style testing indicated compatibility
329 across the tower network of 0.6 ppb CH₄ (Richardson et al., 2017). In this study we use hourly
330 means of CH₄.

331

332 **2.3 Meteorological data**

333 Wind data were measured at the Indianapolis International Airport (KIND), Eagle Creek Airpark
334 (KEYE), and Shelbyville Municipal Airport (KGEZ). The data used are hourly values from the
335 Integrated Surface Dataset (ISD) (<https://www.ncdc.noaa.gov/isd>) and 5-minute values directly
336 from the Automated Surface Observing System (ASOS). A complete description of ASOS
337 stations is available at <https://www.weather.gov/media/asos/aum-toc.pdf>. The accuracy of the
338 wind speed measurements are ± 1 m/s or 5% (whichever is greater) and the accuracy of the wind
339 direction is 5 degrees when the wind speed is ≥ 2.6 m/s. The anemometers are located at about
340 10 meters AGL. The wind data reported in ISD are given for a single point in time recorded
341 within the last 10 minutes of an hour and are closest to the value at the top of the hour.

342 The planetary boundary layer height (BLH) was determined from a Doppler lidar
343 deployed in Lawrence, Indiana about 15 km to the northeast of downtown. The lidar is a Halo
344 Streamline unit, which was upgraded to have extended range capabilities in January 2016. The
345 lidar continuously performs a sequence of conical, vertical-slice, and staring scans to measure
346 profiles of the mean wind, turbulence, and relative aerosol backscatter. All of these
347 measurements are combined using a fuzzy-logic technique to automatically determine the BLH
348 continuously every 20-min (Bonin et al., 2018). The BLH is primarily determined from the
349 turbulence measurements, but the wind and aerosol profiles are also used to refine the BLH
350 estimate. The BLHs are assigned quality-control flags that can be used to identify times when
351 the determined BLH is unreliable, such as when the air is exceptionally clean, the BLH is below
352 a minimum detectable height, or clouds and fog that attenuate the lidar signal exist. Additional
353 details about the algorithm and the lidar operation for the INFLUX project are provided in Bonin

354 et al. (2018). Doppler lidar measurements are available at
355 <https://www.esrl.noaa.gov/csd/projects/influx/>.

356

357 **2.4 Urban methane background**

358 Both aircraft mass balance and inverse modeling methodologies rely on an accurate estimation of
359 the urban CH₄ enhancement relative to the urban CH₄ background in order to produce a reliable
360 flux estimate (Cambaliza et al., 2014; Lamb et al., 2016). The CH₄ mole fraction enhancement is
361 defined as,

$$C_{enhancement} = C_{downwind} - C_{bg} \quad (1)$$

362 where $C_{downwind}$ is the CH₄ mole fraction measured downwind of a source and C_{bg} is the CH₄
363 background mole fraction, which can be measured upwind of the source, but this is not
364 necessary. Background, as defined in this body of literature, is a mole fraction measurement that
365 does not contain the influence of the source of interest, but which is assumed to accurately
366 represent mole fractions that are upwind of the source of interest and measured simultaneously
367 with the downwind mole fractions.

368 Aircraft mass balance studies of Indianapolis mentioned used two main methods to
369 determine a background value. The first method calculates an average of the aircraft transect
370 edges that lie outside of the city domain (Cambaliza et al., 2014). In the second approach, a
371 horizontally varying background is introduced by linearly interpolating median background
372 values of each of the transect edges (Heimbürger et al., 2017). In theory there is also a third
373 method that uses an upwind transect as a background field, but in the studies above it was
374 assumed that the edges are representative of an upwind flow. In the case of an inversion, it is
375 common to pick a tower that is located generally away from urban sources and has on average

376 the smallest overall enhancement (Lavaux et al., 2016). Because choosing the background
377 involves a degree of subjectivity (Heimburger et al., 2017) we consider how these choices may
378 influence emission estimates and introduce error, both random and systematic, using data from
379 the INFLUX tower network.

380 Using tower network data from November 2014 through the end of 2016, two CH₄
381 background fields are generated for the city of Indianapolis based on two different sets of
382 criteria. The notion is based on the fact that a choice of background is currently rather arbitrary
383 in the literature (Heimburger et al., 2017) and at every point in time it is possible to choose
384 multiple background values that are equally acceptable for the flux estimation. In our case both
385 approaches identify a tower suitable to serve as a background for each of the eight wind
386 directions (N, NE, E, SE, S, SW, W, NW), where an arc of 45° represents a direction (e.g. winds
387 from N are between 337.5° and 22.5°). Estimating background for different wind directions is
388 implemented to more accurately represent upwind flow that is hopefully not contaminated by
389 local sources.

390 Criterion 1 corresponds to a typical choice of a background in a case of tower inversion
391 and is based on the concept that the lowest CH₄ mole fraction measured at any given time is not
392 affected by the city sources and therefore is a viable approximation of the background CH₄ mole
393 fractions outside of the city (Miles et al., 2017; Lauvaux et al., 2016). Given this assumption, the
394 tower with the lowest median of the CH₄ enhancement distribution (calculated by assuming the
395 lowest measurement among all towers at a given hour as a background) for each of the wind
396 directions over the November 2014 through December 2016 time period is chosen as a
397 background site (Miles et al., 2017). Criterion 2 requires that the tower is outside of Marion
398 County (outside of the city boundaries) and is not downwind of any known regional CH₄ source

399 (Fig. 2). For some wind directions, there are multiple towers that could qualify as a background;
400 we pick towers in such a manner that they are different for each criterion given a wind direction
401 in order to calculate the error associated with the use of different but acceptable backgrounds.
402 The towers used for both criteria and for each of the eight wind directions are displayed in Table
403 1. Quantifying differences between these two backgrounds allows for an opportunity to better
404 understand the degree of uncertainty that exists in the atmospheric CH₄ background at
405 Indianapolis.

406 To make the comparison as uniform as possible only data from 12-16 LST are utilized
407 (all hours are inclusive) when the boundary layer is typically well-mixed (Bakwin et al., 1998).
408 A lag 1 autocorrelation is found between 12-16 LST hours, i.e. the hourly afternoon data are
409 correlated to the next hour, but the correlation is not significant for samples separated by two
410 hours or more. Therefore, hours 13 and 15 LST are eliminated to satisfy the independence
411 assumption for hourly samples. Furthermore, we make an assumption that the data satisfy steady
412 state conditions. If the difference between consecutive hourly wind directions exceeds 30
413 degrees or the difference between hours 16 and 12 LST exceeds 40 degrees, the day is
414 eliminated. Days with average wind speeds below 2 m/s are also eliminated due to slow
415 transport across the city (the transit time from tower 1 to tower 8 is about 7 hours at a wind speed
416 of 2 m/s).

417

418 **2.5 Frequency and bivariate polar plots**

419 Frequency and bivariate polar plots are used in this work to gain more knowledge regarding CH₄
420 background variability based on criteria 1 and 2, and to identify sources located within the city.

421 To generate these polar plots, we use the *openair* package (from R programming language)

422 created specifically for air quality data analysis (Carslaw and Ropkins, 2012). Bivariate and
423 frequency polar plots indicate the variability of a pollutant concentration at a receptor (such as an
424 observational tower) as a function of wind speed and wind direction, preferably measured at the
425 location of the receptor or within several kilometers of the receptor. The frequency polar plot is
426 generated by partitioning the CH₄ hourly data into the wind speed and direction bins of 1 m s⁻¹
427 and 10° respectively. To generate bivariate polar plots, wind components u and v are calculated
428 for hourly CH₄ mole fraction values, which are fitted to a surface using a Generalized Additive
429 Model (GAM) framework in the following way,

$$\sqrt{C} = \beta + s(u, v) + \epsilon \quad (2)$$

430 where C is the CH₄ mole fraction transformed by a square root to improve model diagnostics
431 such as a distribution of residuals, β is mean of the response, s is the isotropic smoothing
432 function of the wind components u and v , and ϵ is the residual. For more details on the model
433 see Carslaw and Beevers (2013).

434

435 **2.6 Temporal variability and approximate flux estimation**

436 Temporal variability may play an important role in the quantification of urban CH₄ emissions.
437 Lamb et al. (2016) suggested that temporal variability might partially explain the differences
438 among CH₄ flux estimates shown in Figure 1. If temporal variability of CH₄ emissions exists
439 within the city, disagreements in the CH₄ flux between studies could be attributed to differences
440 in their sampling period. Because the INFLUX tower data at Indianapolis contain measurements
441 at all hours of the day over multiple years, we can utilize this dataset to better understand the
442 temporal variability in methane emissions in the city.

443 We apply a simplified atmospheric boundary layer budget, not to estimate precisely the
 444 actual city emissions, but rather to evaluate temporal variability of the emissions. We begin by
 445 assuming CH₄ emissions Q_a (mass per unit time per unit area) are not chemically active and are
 446 constant over a distance Δx spanning a significant portion of the city. The next assumption is
 447 that a CH₄ plume measured downwind of the city is well mixed within a layer of depth H (which
 448 is the same as BLH). We treat wind speed u as constant within the layer for every hour
 449 considered. Given the above-mentioned assumptions we can write a continuity equation
 450 describing mass conservation of CH₄ concentration C within a box in the following fashion,

$$\Delta x H \frac{\partial C}{\partial t} = \Delta x Q_a + uH(C_b - C) + \Delta x \frac{\partial H}{\partial t} (C_a - C) \quad (3)$$

451 where C_b is the CH₄ concentration upwind of the city (or background), and C_a is the CH₄
 452 concentration above the mixed layer (Hanna et al., 1982; Arya, 1999; Hiller et al., 2014). The
 453 left hand side of the equation represents the change in CH₄ concentration with time, $\Delta x Q_a$
 454 denotes a constant CH₄ source over the distance Δx , $uH(C_b - C)$ indicates a change of CH₄
 455 concentration due to horizontal advection, and finally $\Delta x \frac{\partial H}{\partial t} (C_a - C)$ term accounts for the
 456 vertical advection and encroachment processes that result from changing BLH. By assuming
 457 steady state conditions ($\frac{\partial C}{\partial t} = 0$ and $\frac{\partial H}{\partial t} = 0$), the equation can be simplified to

$$Q_a = \frac{uH(C - C_b)}{\Delta x} \quad (4)$$

458 We use equation 4 to estimate hourly CH₄ emissions (Q_a) from Indianapolis (see
 459 assumptions in the paragraph below) given hourly averaged data of H from the lidar positioned
 460 in the city, wind speed (u) from the local weather stations, and upwind (C_b) and downwind (C)
 461 CH₄ mole fractions measured (and then converted to concentrations) at towers 1, 8, and 13

462 (depending on a wind direction) using data from heights of 40 m, 41 m, and 87 m respectively
463 (see Fig. 2).

464 The CH₄ concentrations are derived from CH₄ mole fractions by approximating average
465 molar density of dry air (in mol m⁻³) within the boundary layer for every hour of the day, where
466 variability of pressure with altitude is calculated using barometric formula and it is assumed that
467 temperature decreases with altitude by 6.5 K per kilometer. The hourly surface data for pressure
468 and temperature are taken from KIND weather station. The difference between concentrations
469 $C - C_b$ is instantaneous and not lagged, where C_b represents air parcel entering the city and C
470 represents the same air parcel exiting the city (Turnbull et al., 2015). The CH₄ enhancements
471 $C - C_b$ are estimated for daytime by averaging observations spanning 12-16 LST and for
472 nighttime by averaging observations spanning 20-5 LST. These time periods are based on lidar
473 estimations of when on average H varies the least. The day and night were required to contain at
474 least 3 and 9 hourly CH₄ values respectively for averaging to occur, otherwise the day/night is
475 eliminated. Observations when H is below 100 m are not used to avoid the cases when
476 measurements from towers may be above the boundary layer. In order to better achieve the
477 assumption that the boundary layer is fully mixed (especially at night), all hours with wind
478 speeds below 4 m/s are eliminated (Van De Wiel., 2012). To approximate the emissions of the
479 whole city we need to know the approximate area of the city and the distance over which the
480 plume is affected by the city CH₄ sources. The area of the city is about 1024 km² (the area of
481 Marion County) and the length that plume traverses when it is over the city ranges from 32 to 35
482 km depending on which downwind tower is used. We assume that CH₄ measurements at towers
483 8 and 13 are representative of a vertically well-mixed city plume as the towers are located
484 outside of the city boundaries and allow for sufficient vertical mixing to occur. For S and SW

485 wind directions tower 8 observations are used to represent downwind conditions with
486 background observations coming from towers 1 and 13, respectively (based on criterion 1 shown
487 in Table 1). For W wind direction, tower 13 observations represent the downwind with
488 background obtained from tower 1. The wind direction is required to be sustained for at least 2
489 hours, otherwise the data point is eliminated.

490

491 **2.7 Indianapolis CH₄ sources**

492 Only a few known CH₄ point sources exist within Indianapolis (Cambaliza et al., 2015, Lamb et
493 al., 2016). The Southside Landfill (SSLF), located near the center of the city, is thought to be the
494 largest point source in the city with emissions ranging between about 28 mol/s (inventory from
495 Maasackers et al. (2016), GHG reporting program, and inverse estimates from ground-based
496 mobile sampling employed in Lamb et al. (2016)) and 45 mol/s (aircraft; Cambaliza et al.
497 (2015)) depending on an emission estimation methodology. However, using Cambaliza et al.
498 (2015) aircraft data and applying a different background formulation Lamb et al. (2016) found
499 emission values of SSLF closely agreeing with 28 mol/s estimate. SSLF could account for as
500 little as 33% (top-down from Cambaliza et al., 2015) or as much as 63% (inventory from
501 Maasackers et al., 2016) of total Marion County CH₄ emissions. Other city point sources are
502 comparatively small; the wastewater treatment facility located near SSLF contributes about 3-7
503 mol/s (inventory from Lamb et al. 2016), and the transmission-distribution transfer station at
504 Panhandle Eastern Pipeline (also known as a city gate and further in this study abbreviated as
505 PEP) is estimated to be about 1 mol/s (inventory from Lamb et al. 2016). The remaining CH₄
506 sources, mainly from NG infrastructure leaks and livestock, are considered to be diffuse sources
507 and are not well known. Potential sources of emissions related to NG activities include gas

508 regulation meters, transmission and storage, distribution leaks, and Compressed Natural Gas
509 (CNG) fleets. These diffuse NG sources account for 21-67% of the city emissions or 20 mol/s
510 (inventory from Maasakkers et al., 2016) to 64 mol/s (top down from Cambaliza et al., 2015).
511 Livestock emissions for Marion County are estimated to be around 1.5 mol/s (inventory from
512 Maasakkers et al., 2016). These prior studies present conflicting conclusions regarding the
513 magnitude of the diffuse NG CH₄ source in Indianapolis.

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515 3 Results and discussion

517 3.1 Inversion and city boundaries

518 A significant portion of CH₄ emissions across the U.S. can be characterized by numerous
519 relatively large point sources scattered throughout the country rather than by broad areas of
520 smaller enhancements (Maasakkers et al., 2016). Because of this, the total emissions for a given
521 domain can be very sensitive to how that domain is defined. A small increase or decrease in the
522 domain area could add or remove a large point source and significantly impact the total
523 emissions defined within the domain.

524 In the case of Indianapolis, this issue became apparent when the emissions were
525 calculated using an atmospheric inversion model (Lamb et al., 2016; Lauvaux et al., 2016). The
526 atmospheric inversion solved for fluxes in domain 1 (Fig. 2), which significantly increased the
527 estimated emissions in comparison with the inventory values that were gathered mainly within
528 Marion County (domain 2). When reduced to domain 2, inverse modeling emission estimate
529 decreases to 107 mol/s (from about 160 mol/s), which falls within an error bar of Lamb et al.
530 (2016) inventory estimate. This difference is significant and could at least partially explain the
531 discrepancy shown in Figure 1 between the emission values from the inventories and emission

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541 results from the inverse modeling. However, even the decreased inverse modeling estimate is
542 about 91% higher than the inventories.

543 Additionally, the subject of the domain is relevant for airborne mass balance flights
544 because a priori the magnitude and variability of background plume is unknown and could be
545 easily influenced by upwind sources. The issue of background is discussed further in the next
546 section.

547

548 3.2 Variability in CH₄ background

549 Comparisons between criterion 1 and criterion 2 CH₄ background mole fractions as a
550 function of wind speed and direction are visualized using frequency and bivariate polar plots
551 (Fig. 4). Both backgrounds generally agree on the higher CH₄ originating from the SW, SE, and
552 E wind directions (Figs. 4c-f); however, the values themselves differ especially when winds are
553 from NW, SW, and SE. As background difference plots (Figs. 4g-h) indicate, there is a
554 noticeable variability between the magnitudes of the CH₄ backgrounds, where criterion 2, by
555 design, typically has higher background mole fractions. The background differences, at a given
556 hour, suggest that the CH₄ field flowing into the city is heterogeneous with differences between
557 towers ranging from 0 to over 45 ppb (Fig. 4g). Because large gradients in CH₄ background over
558 the city could pose challenges for flux estimations using top down methods such as inverse
559 modeling and aircraft mass balance, it is imperative to establish whether the background
560 differences vary randomly or systematically and how to choose a background to minimize these
561 errors.

562 To further understand the nature of background variability we calculate the mean,
563 standard deviation, and standard error of background hourly differences between criterion 2 and

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565 criterion 1 from November 2014 to December 2016 for each of the eight wind directions
566 mentioned in Table 1. The results are shown in Figure 5. Systematic bias is evident for the SE,
567 S, SW, W, and NW wind sectors, whereas random error dominates N, NE, and E wind
568 directions. Wind directions showing statistically significant bias have mean biases ranging from
569 2 to 5 ppb, with values as large as 8 ppb falling within the range of $2 \times$ standard error. Standard
570 deviation plot indicates potential background discrepancy that can occur on any given day, where
571 W wind direction is the least variable with $2 \times$ standard deviation close to 20 ppb, while SE wind
572 direction is the most variable with $2 \times$ standard deviation falling at about 50 ppb.

573 Random errors in the mole fractions of background differences (biases) are also
574 important and are a function of the length of the data record. We quantify the random error in
575 the CH₄ background mole fraction differences using the bootstrap method by randomly sampling
576 2 to 150 hours (small and large sample size) of the background CH₄ differences for each of the
577 wind directions with replacement (we make the assumption that our differences are independent
578 since we eliminated lag 1 autocorrelation from the data). This sub-sampling experiment is
579 repeated 5000 times (Efron and Tibshirani, 1986). The standard deviations of the mean
580 (standard error) of the 5000 simulated differences are calculated for each wind direction. The
581 resulting standard errors of the city CH₄ background differences, multiplied by 2 to represent the
582 95% confidence intervals, are shown as a function of the length of the data record in Figure 6.
583 Because random error falls as sample size grows it makes sense to assign a threshold indicating a
584 minimum number of samples needed to achieve a theoretical precision for each wind direction.

585 One way to assign a required precision would be to make sure that the standard error
586 (random error) reaches a point where it is less than Indianapolis enhancement of about 12 ppb (a
587 higher estimate of the Indianapolis enhancement from section 3.3) by a factor of 2 when

588 combined with a bias (Table 2). Meaning that the sum of bias and standard error must be at most
589 6 ppb. In this approach each wind direction would have a different threshold because of the
590 differences in biases. For instance, given this requirement NW direction would need a random
591 error of 1 since its bias is 5. For NW direction, this threshold would require more than 150
592 samples. For N direction on the other hand, where the bias is 1, the requirement is fulfilled when
593 random error crosses 5 ppb at 74 samples. Now we consider these random and systematic errors
594 in CH₄ background differences in the context of Indianapolis urban CH₄ emissions.

595 For Indianapolis, using INFLUX tower network, we estimated that depending on sample
596 size (number of hours sampled) and wind direction, background gradient across the city over 12-
597 16 LST could vary from 0 to about 50 ppb (Fig. 5b). Given that the average afternoon CH₄
598 enhancement of the city is around 8-12 ppb (section 3.3; Fig. 7; Cambaliza et al., 2015; Miles et
599 al., 2017), the error on the estimated emissions could easily be over 100% if the analysis does not
600 approach the issue of background with enough sampling. A sample size of about 50 independent
601 hours significantly decreases background uncertainty for N, NE, E, S, and W wind directions and
602 allows for a more accurate assessment of the CH₄ emissions at Indianapolis. For CH₄ sources
603 with a significantly larger signal than their regional background, the mentioned background
604 variability becomes less impactful on results, but because Indianapolis is a relatively small
605 emitter of CH₄, and because there are relatively large sources outside of the city, uncertainties
606 due to background estimation are comparatively large. Our uncertainty assessment suggests that
607 the highly variable CH₄ emission values of Indianapolis from aircraft mass balance calculations
608 shown in Figure 1 are at least partially due to the variability in the urban CH₄ background of
609 Indianapolis.

610

611 3.3. Temporal variability of methane enhancements and fluxes in Indianapolis

612 Figure 7 presents average CH₄ mole fraction enhancements and flux calculations
613 (equation 4) at towers 8 and 13 for years 2014, 2016, and 2013-2016 (for the detailed
614 methodology see section 2.6). The years of 2014 and 2016 are chosen for temporal comparison
615 because they do not contain major BLH data gaps. The error bars in the figure show the standard
616 error multiplied by 2 indicating 95% confidence interval of each average.

617 One of the more interesting features in the Figure 7 is a day/night variability of CH₄
618 emissions at Indianapolis. The most prominent example of this feature is found in Figure 7c,
619 where the estimates for both years suggest that daytime emissions are approximately twice as
620 large as the emissions at night. The decrease of the CH₄ emissions at night also appears in tower
621 13, but the errors are too high in those estimates to make any definitive conclusions. A similar
622 urban CH₄ emissions diurnal variability is reported by Helfter et al. (2016) in their study of
623 GHGs for London, UK, where they attribute diurnal variation of CH₄ emissions to the NG
624 distribution network activities, fugitive emissions from NG appliances, and to temperature-
625 sensitive CH₄ emission sources of biogenic origin (such as a landfill). Taylor et al. (2018)
626 suggest that CH₄ emissions from landfills exhibit a diurnal cycle with higher emissions in early
627 afternoon and 30-40% lower emissions at night.

628 With regard to yearly temporal variability we are only able to compare years 2014 and
629 2016 due to limited BLH data for other years. Results from both towers suggest that
630 Indianapolis overall CH₄ emissions did not change significantly between 2014 and 2016.
631 Although it is important to be cautious about interpreting actual flux estimations given the
632 assumptions mentioned in section 2.6, it is interesting to note that the flux values from both
633 towers average to about 70 mol/s, which puts our value right in between inventory and inversion

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635 estimates shown in Figure 1. If we assume that SSLF emissions are generally known (GHG
636 reporting program) that would indicate that emissions from NG distribution are likely to be **about**
637 **14 mol/s (70%)** higher than **what** both of the inventories currently estimate **but within the error**
638 **bars of** Lamb et al. (2016)'s inventory calculation. Another possible scenario is that SSLF
639 emissions are higher than what is currently assumed. Given these complexities, uncertainty
640 regarding the exact emissions from NG distribution at Indianapolis still remains.

641

642 3.4 Methane Sources in Indianapolis

643 Bottom-up emission inventories have difficulty tracking changes in sources over time. Our
644 continuous tower network observations can monitor temporal and spatial variability in sources of
645 CH₄ in Indianapolis. To do so we employ the aforementioned bivariate polar plots to verify
646 known sources and potentially identify unknown sources across the city. We compare two time
647 periods, 2014-2015 (two full years) and 2016. Figure 8 displays bivariate polar plots of CH₄
648 enhancements using criterion 1 background at 9 INFLUX towers in Indianapolis over the two
649 years of 2014 and 2015. Figure 9 shows the same plot, but for the year 2016. Here we have
650 separated 2016 from 2014-2015 because of different results noted during these times.

651 The images reveal that the most consistent and strongest source in the city is the SSLF.
652 This is most evident from the 40+ ppb CH₄ enhancements detected at towers 7, 10 and 11
653 coming from the location of the SSLF (by triangulation). Enhancements from the landfill appear
654 to also be detectable at towers 2, 4, 5, and 13. Based on these observations it can be concluded
655 that there are no other point sources in Marion County comparable in size to the SSLF. A small
656 fraction of the SSLF plume is likely due to the co-located wastewater facility, but the inventory
657 estimates suggest that the wastewater treatment facility is responsible for no more than 7% of

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662 this plume (Cambaliza et al., 2015; Massackers et al., 2016). The PEP, located in the
663 northwestern section of the city, may be partially responsible for a plume of 5-10 ppb at towers 5
664 and 11. However, the plume is less detectable using the criterion 2 background value that has
665 higher background (using tower 8 as a background) from NW wind direction (not shown),
666 adding uncertainty to the true magnitude of the enhancement from this source. The same is true
667 for towers 2 and 13, which have pronounced plumes when winds are from the NW with the
668 criterion 1 background, but when background 2 is used these plumes vanish (not shown). Such
669 inconsistency makes it difficult to attribute these plumes to a specific source.

670 Another important point is the cluster of large enhancements surrounding tower 10 in
671 2014-2015. Because no other tower sees these enhancements (at least at comparable
672 magnitudes), we believe that they are the result of nearby NG leaks. These plumes are not
673 consistent temporally or spatially as they mostly disappear in 2016, potentially indicating that
674 they are transient and localized NG distribution leaks. It is difficult to ascertain the exact
675 combined magnitude of these leaks since they mix together with SSLF into an aggregated city
676 plume when observed from downwind towers such as 8 and 13. None of the individual leaks
677 appears to be similar in magnitude to the emissions that originate from SSLF. Diffuse NG
678 emissions comparable to the SSLF source (Lamb et al., 2016) may exist. Our flux estimations at
679 towers 8 and 13, however, imply that the magnitude of NG diffuse source suggested by the top-
680 down analyses in Cambaliza et al. (2015) and Lamb et al. (2016) are probably overestimates (see
681 section 3.3). We hypothesize that the relatively high Indianapolis CH₄ emissions (see Fig. 1)
682 reported by Cambaliza et al. (2015) could be a result of random errors in upwind conditions (see
683 section 3.2) influencing the small number of airborne flux estimates.

684

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755 **4 Conclusions**

756 We have examined four potential contributions to discrepancies between urban top-down and
757 bottom-up estimates of CH₄ emissions from Indianapolis: domain definition, heterogeneous
758 background mole fractions, temporal variability in emissions, and sources missing from
759 inventories. Results indicate that the urban domain definition is crucial for the comparison of the
760 emission estimates among various methods. Atmospheric inverse flux estimates for Marion
761 County, which is similar to the domain that is analyzed by inventory and airborne mass balance
762 methodologies (Mays et al., 2009, Cambaliza et al., 2014, Lamb et al., 2016), is 107 mol/s
763 compared to 160 mol/s that is estimated for the larger domain (Hestia inventory domain; Gurney
764 et al., 2012). This partially explains higher emissions in inverse modeling estimates shown by
765 Lamb et al. (2016); however, 107 mol/s is still 91% higher than what EPA and Lamb et al.
766 (2016) find in their inventories (Fig. 1).

767 To better understand background variability at Indianapolis two different but acceptable
768 background estimates, based on specific criteria for each wind direction, and their differences are
769 used to assess heterogeneity of CH₄ background at Indianapolis. Background criterion 1 looks
770 for a tower that is consistently lower than other towers, while background criterion 2 picks a
771 tower that is outside of Marion County domain and is not downwind of any nearby sources as
772 determined by EPA 2012 inventory. We focus on midday atmospheric conditions to avoid the
773 complexities of vertical stratification in the stable boundary layer. The midday Indianapolis
774 atmospheric CH₄ mole fraction background is shown to be heterogeneous with 2-5 ppb
775 statistically significant biases for NW, W, SW, S and SE wind directions. Random errors of
776 background differences are a function of sample size and decrease as a number of independent
777 samples increase. Small sample sizes, such as a few hours of data from a single point, are prone

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779 to random errors on the order of 10-30 ppb in the CH₄ background, similar to the magnitude of
780 the total enhancement from the city of Indianapolis, which is estimated to be on average around
781 10-12 ppb. Longer-term sampling and/or more extensive background sampling are necessary to
782 reduce the random errors. Sample size required to reduce random errors of background
783 differences to an acceptable value for flux calculation is largely dependent on a wind direction.
784 Both bias (long-term average of background differences) and its random error are important
785 when estimating total background uncertainty. The results indicate that N, NE, E, S, and W
786 wind directions are more favorable for flux estimation and would require multiple days of
787 measurements (e.g. about 50 independent hours of measurements) to reduce background
788 uncertainty to about 6 ppb, which is half the magnitude of the typical CH₄ enhancement from
789 Indianapolis. The remaining wind directions would require over 150 independent hourly
790 measurements to achieve similar precision. We also estimate that depending on a wind direction
791 for any given hour the spatial variability in background can be anywhere from 0 to 50 ppb. This
792 uncertainty in the CH₄ background may partially explain Heimbürger et al. (2017) finding of
793 large variability in airborne estimates of Indianapolis CH₄ emissions. Given many samples, the
794 airborne studies converge to an average value of CH₄ flux that is noticeably closer to the
795 inventory estimates for Indianapolis than several of the individual estimates presented in Figure
796 1.

797 Measurement and analysis strategies can minimize the impacts of these sources of error.
798 Spatially extensive measurement of upwind CH₄ mole fractions are recommended. For towers or
799 other point-based measurements, multiple upwind measurement locations are clearly beneficial.
800 For the aircraft mass balance approach, we recommend an upwind transect to be measured,
801 lagged in time if possible, to provide a more complete understanding of the urban background

802 conditions. Complex background conditions might suggest that data from certain days or wind
803 directions should not be used for flux calculation. Finally, a mesoscale atmospheric modeling
804 system informed with the locations of important upwind CH₄ sources can serve as a powerful
805 complement to the atmospheric data (Barkley et al., 2017). Such simulations can guide sampling
806 strategies, and aid in interpretation of data collected with moderately complex background
807 conditions.

808 With regard to temporal variability, no statistically detectable changes in the emission
809 rates were observed when comparing 2014 and 2016 CH₄ emissions. However, a large
810 difference between day and night CH₄ emissions was implied from a simple budget estimate.
811 Night (20-5 LST) emissions may be 2 times lower than the emissions during the afternoon (12-
812 16 LST) hours. Because prior estimates of top-down citywide emissions are derived using
813 afternoon-only measurements, overall emissions of Indianapolis may be lower than these studies
814 suggest. This bias may be present in studies performed in other cities as well. Our study
815 suggests that day/night differences in CH₄ emissions must be understood if regional emission
816 estimates are to be calculated correctly. Long-term, tower-based observations are an effective
817 tool for understanding and quantifying multi-year variability in urban emissions.

818 One final point addressed in this study is the location of major CH₄ sources in
819 Indianapolis. Analysis of the INFLUX tower observations suggest a diffuse NG source that
820 exceeds both of the inventory estimates by 70%, but additionally our analysis shows that the
821 discrepancy is less than that proposed by highest values reported in Lamb et al. (2016) (see Fig.
822 1). Uncertainty remains regarding the magnitude of the diffuse NG source of CH₄. The only
823 major point source in the city is SSLF and it is observed at multiple towers. There is an evidence

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836 | for occasional point-source NG leaks, but they appear to be transient in time, and limited in their
837 | strength.

838 | Overall, assessment of the CH₄ emissions at Indianapolis highlights a number of
839 | uncertainties that need to be considered in any serious evaluation of urban CH₄ emissions. These
840 | uncertainties amplify for Indianapolis since the enhancement signal from its CH₄ emissions is
841 | comparable in magnitude to variability in the regional background flow and as our results show
842 | it may be difficult at times to distinguish noise in the background from the actual city emissions
843 | signal. The evaluation of larger CH₄ sources may be easier with respect to separating signal
844 | from background. However, all of the points raised in this work will be nonetheless relevant and
845 | need to be addressed for our understanding of urban CH₄ emissions to significantly improve.

846

847 | **Author Contribution**

848 | Nikolay Balashov, Kenneth Davis, and Natasha Miles developed the study and worked together
849 | on generating the main hypothesis of this work. They also wrote most of the manuscript.
850 | Nikolay Balashov wrote all of the codes and performed the analyses presented in this work as
851 | well as generated all of the figures. Natasha Miles and Scott Richardson helped with
852 | maintenance and gathering of the INFLUX tower data. They also wrote section 2.2 of the paper.
853 | Thomas Lauvaux helped with the analysis presented in Fig. 1 and section 3.1 concerning
854 | interpretation of the inversion modeling results from Lamb et al. (2016). Thomas Lauvaux also
855 | helped with repeating the inversion experiment for two different Indianapolis domains (Fig. 1).
856 | Zachary Barkley significantly contributed to discussions regarding the hypothesis and careful
857 | presentation of sections 2.6 and 3.3. Timothy Bonin provided all of the lidar data and wrote the

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863 second part of section 2.3 regarding the lidar and the methodology used to determine planetary
864 boundary layer heights. He also contributed to sections 2.6 and 3.3.

865

866 **Competing Interests**

867 The authors declare that they have no conflict of interest.

868

869 **Acknowledgements**

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871 number 70NANB10H245). We would like to thank Dr. Bram Maasackers for the helpful
872 discussion regarding the EPA 2012 inventory and the relevant error structure. We also thank Dr.
873 Paul Shepson and Dr. Brian Lamb for their useful input regarding airborne mass balance flights
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879 **References**

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 1127 **Tables**

1129 **Table 1.** INFLUX towers used to estimate CH₄ background based on two different criteria. Numbers in
 1130 bold indicate towers chosen to generate a background field when multiple options are possible (for more
 1131 details see discussion). In short, criterion 1 uses towers with the lowest mean CH₄ for a specific wind
 1132 direction, and criterion 2 uses towers outside of Marion County and not downwind of large sources
 1133 (including the city as a whole).

Wind Direction	CH ₄ Background Towers	
	Criterion 1	Criterion 2
North (N)	8	13 , 8
Northeast (NE)	8	13 , 8, 2
East (E)	2 , 8	8 , 4, 1, 2
Southeast (SE)	1	8 , 13, 4, 1
South (S)	1	4 , 13, 1
Southwest (SW)	13	1 , 4
West (W)	1	4 , 1
Northwest (NW)	1	8 , 1

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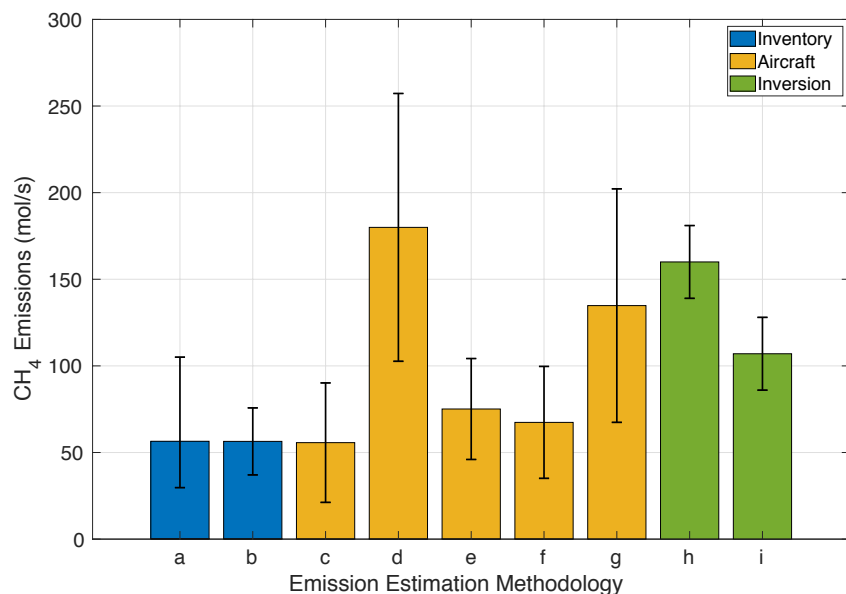
Table 2. A number of independent samples needed (column 4) to satisfy combined requirement of 6 ppb background error based on the sum of bias and random error (explained in section 3.2) as a function of wind direction.

Wind Direction	Bias (ppb)	Threshold (ppb)	Samples Needed
N	1	5	74
NE	1	5	36
E	0.5	5.5	46
SE	4	2	>150
S	2	4	53
SW	4.5	1.5	>150
W	3	3	52
NW	5	1	>150

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Figures

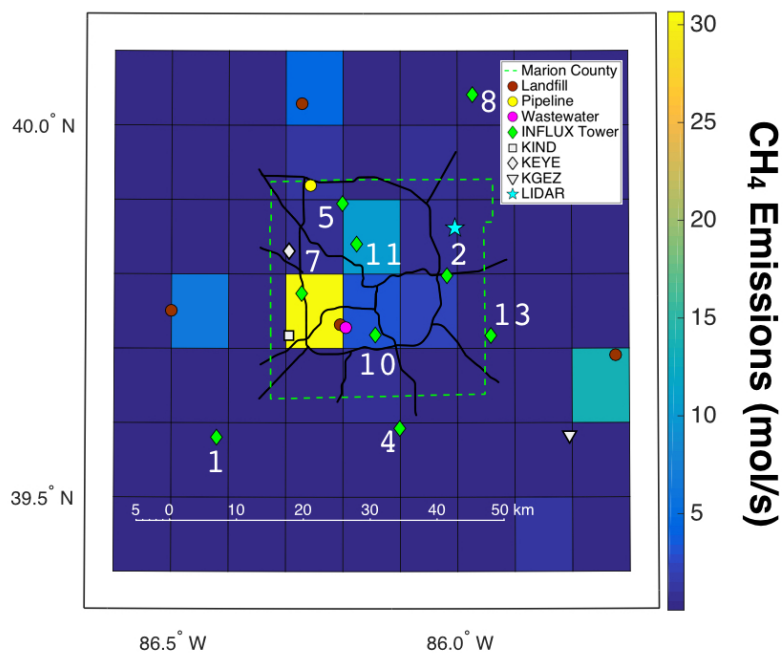


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1189 **Figure 1.** Various estimates of CH₄ emissions at Indianapolis. **(a, b)** Bottom-up estimates of CH₄
1190 emissions conducted by Lamb et al. (2016) in 2013 and Maasackers et al. (2016) based on the EPA 2012
1191 inventory respectively. Error bars show 95% confidence intervals (for more details see above-mentioned
1192 articles). **(c-g)** Top-down evaluations of CH₄ emissions with aircraft from various flight campaigns where
1193 **(c)** contains 5 flights over March-April of 2008, **(d)** contains 3 flights over November-January of 2008-
1194 09, **(e)** contains 5 flights over April-July of 2011, **(f)** contains 9 flights from November-December, 2014,

1195 and (g) contains the same 5 flights over April-July of 2011 as in (e) but uses different methodology.
1196 Methodologies for (c-f) are described in Lamb et al. (2016) and methodology for (g) is described in
1197 Cambaliza et al. (2015). Error bars show 95% confidence intervals (for more details see above-
1198 mentioned articles). (h, i) Top-down evaluations of CH₄ emissions for 2012-2013 using tower inversion
1199 modeling methodology with two different domains, where (h) uses the full domain of Figure 2 and (i)
1200 uses only the Marion County domain of Figure 2. The inversion methodology and 95% confidence
1201 intervals are described in detail in Lamb et al. (2016).

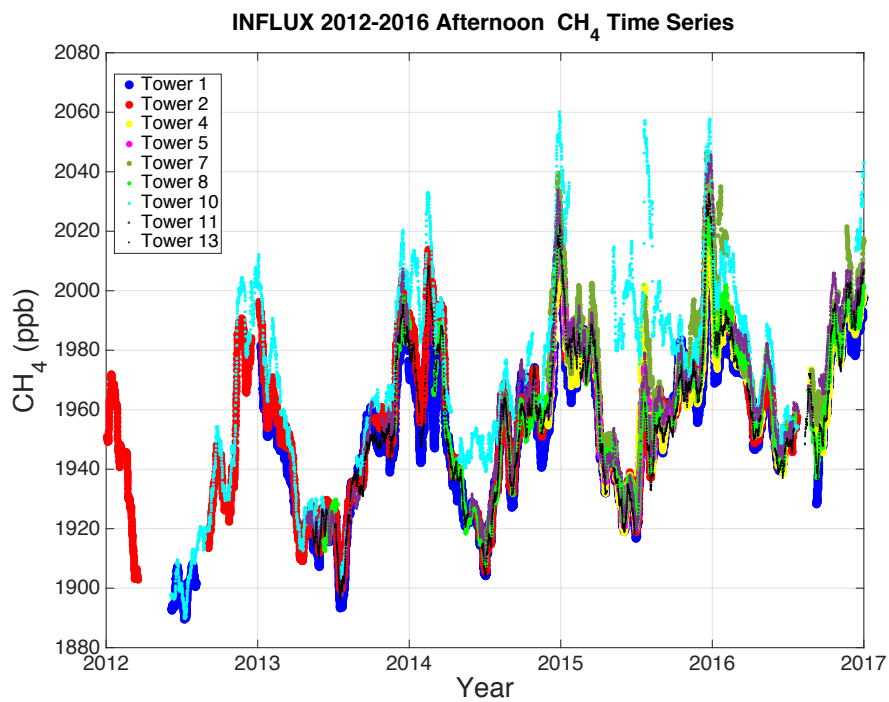
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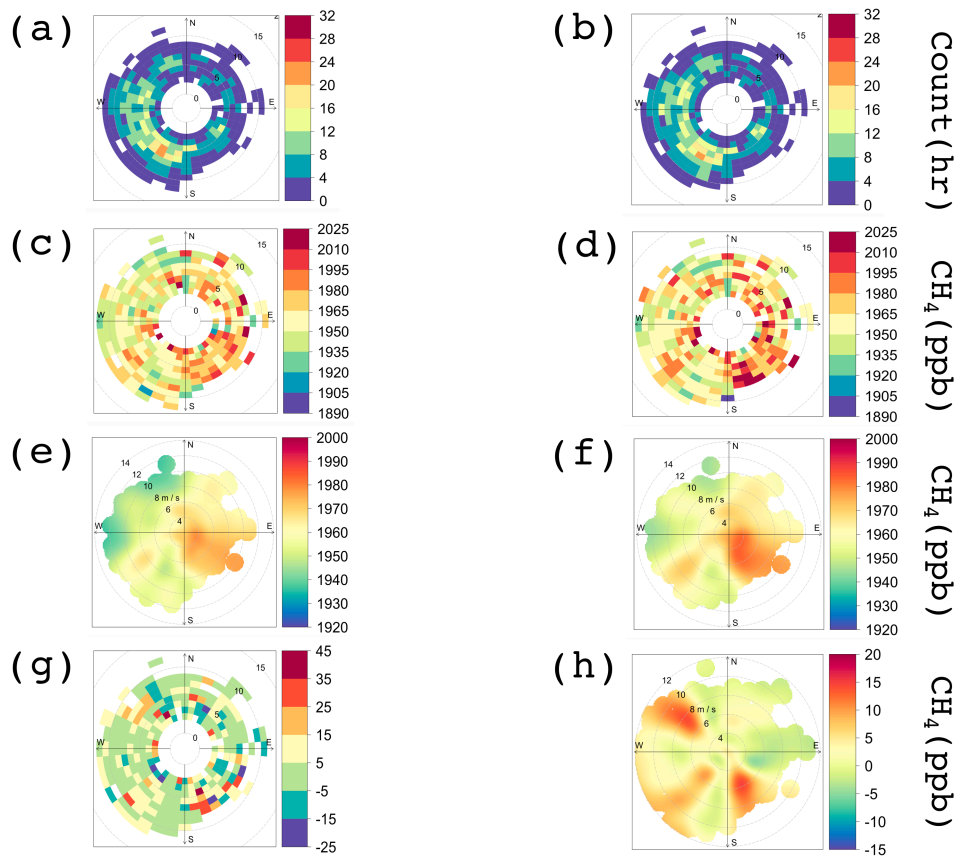
1207 **Figure 2.** Map of the primary roads in Indianapolis, INFLUX towers, lidar system, weather stations, and
1208 a few CH₄ point sources plotted over the gridded CH₄ emissions (mol/s) from the EPA 2012 Inventory
1209 (Maasackers et al., 2016). The gridded map of emissions includes emissions from the mentioned point
1210 sources; their position is provided to aid in interpretation of the observations. The dashed bright green
1211 line denotes Marion County borders.

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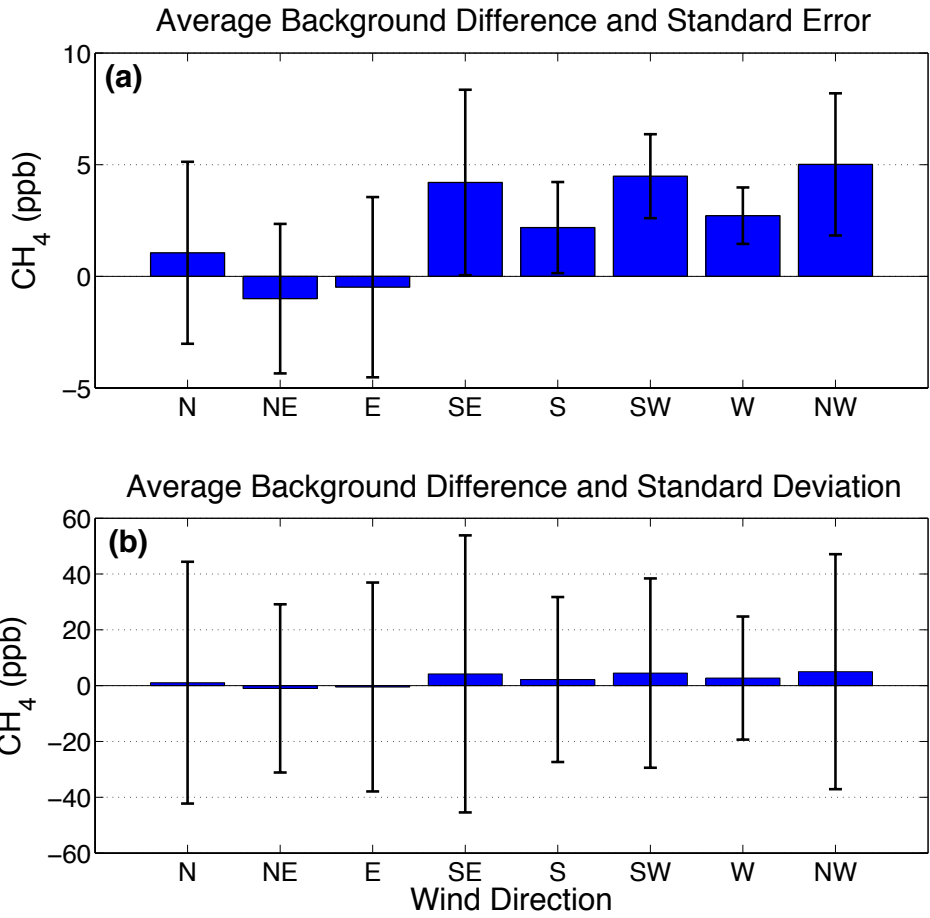
1214 **Figure 3.** 20-day running average of afternoon (12-16 LST; the hours are inclusive) CH₄ mole fractions
1215 as measured by the INFLUX tower network (highest available height is used) from 2012 through 2016.



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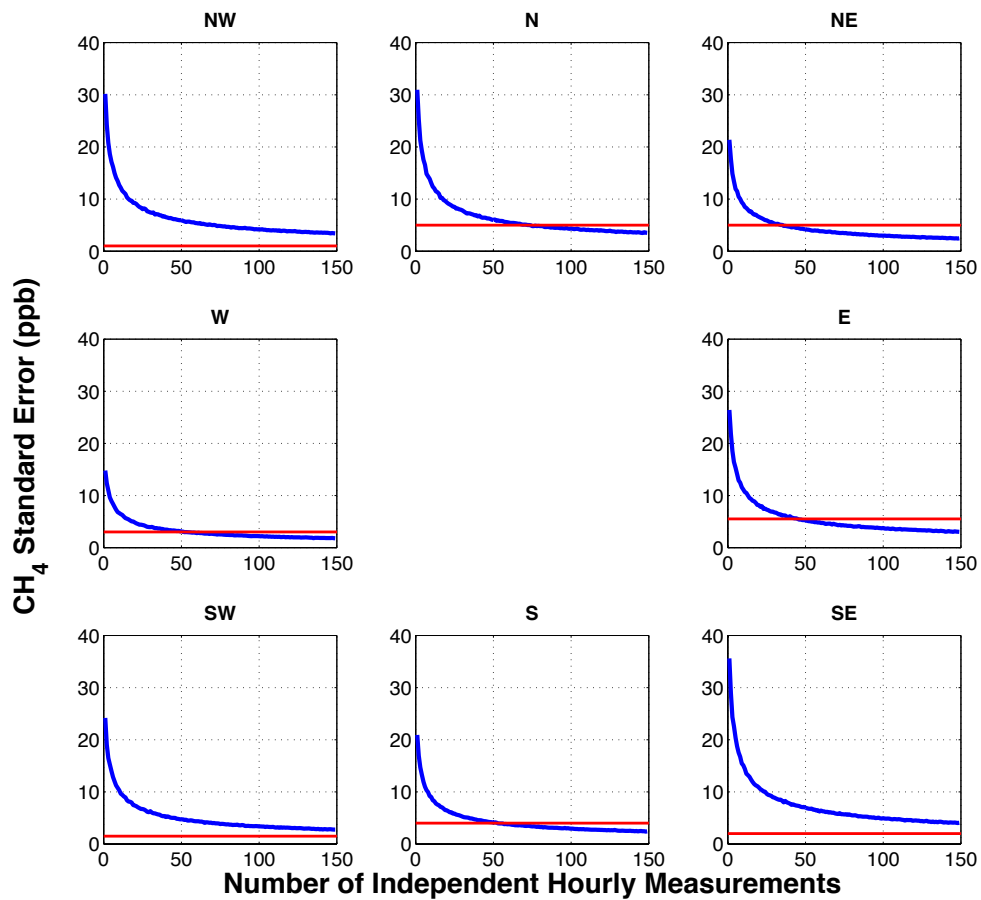
1217 **Figure 4.** Frequency and bivariate polar plots of CH₄ background for Indianapolis using data from 12-16
 1218 LST, November 2014 through December 2016 given 2 different criteria (Table 1). **(a)** Polar histogram
 1219 indicating a number of hourly measurements available using criterion 1. **(b)** Same as (a) only for criterion
 1220 2. Differences between (a) and (b) are due to slight differences in data availability at the considered
 1221 towers. **(c)** Polar frequency plot of the CH₄ background using criterion 1. **(d)** Same as (c) only for
 1222 criterion 2. **(e)** Polar bivariate plot of CH₄ background using criterion 1. **(f)** Same as (e) only for criterion
 1223 2. **(g)** Polar frequency plot of difference between the backgrounds: *criterion 2 – criterion 1*. **(h)** Same
 1224 as (g) but shown with a bivariate polar plot.

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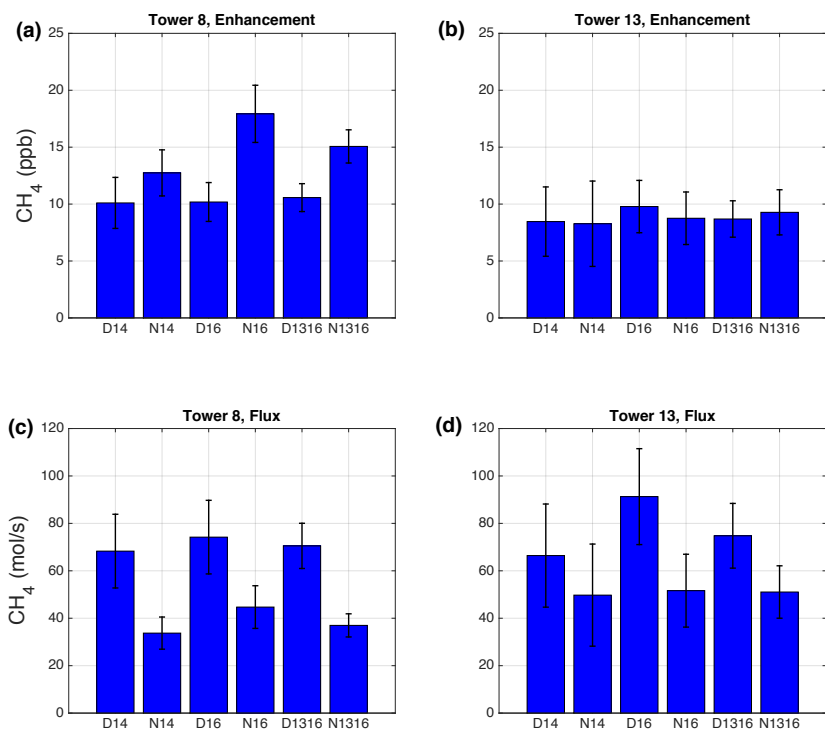
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 1227 **Figure 5.** Average of the differences between criteria 2 and 1 CH₄ backgrounds at Indianapolis as a
 1228 function of wind direction. These averages are generated from the same data that is used in Figure 4 and
 1229 reflect results shown in Figure 4g. Error bars indicate in (a) 2 × standard error and in (b) 2 × standard
 1230 deviation.

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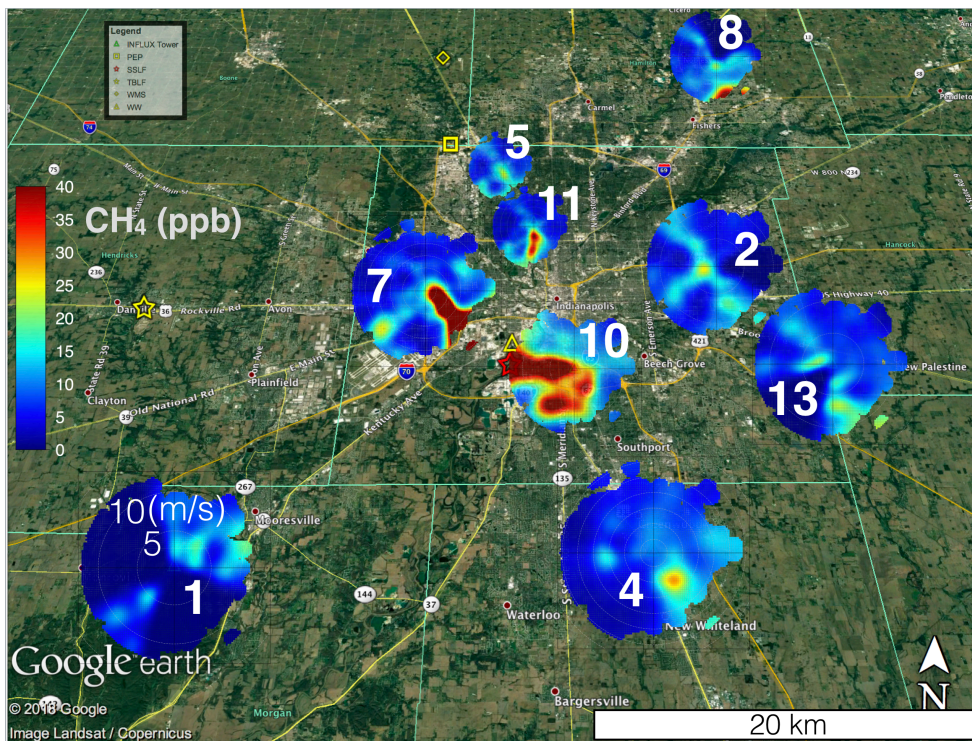
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Figure 6. Bootstrap simulation of the standard errors $\times 2$ in Indianapolis CH₄ background mole fraction differences (between criteria 2 and 1) as a function of sample size and wind direction (see text for details). Thresholds for each of the wind directions indicate a random error threshold needed for the background uncertainty to be within 50% of Indianapolis CH₄ enhancement of 12 ppb.



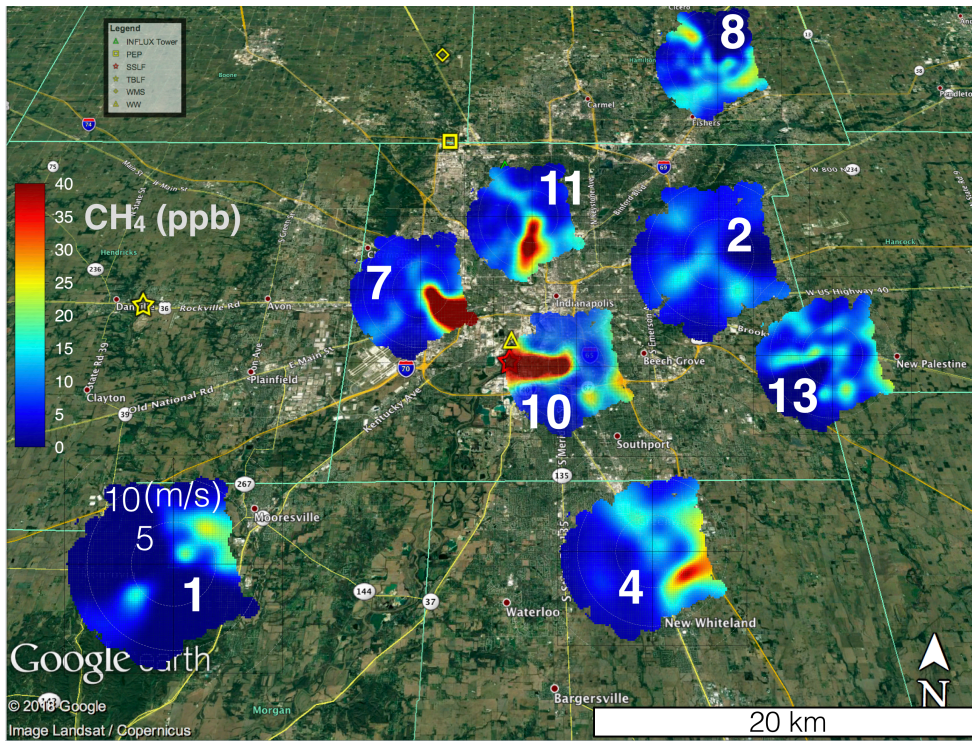
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1239 **Figure 7.** Averages of the daytime (D) and nighttime (N) CH₄ enhancements and fluxes at INFLUX
 1240 towers 8 and 13 for years 2014 (14), 2016 (16), and 2013-2016 (1316). The error bars represent 95%
 1241 confidence interval of each mean value. **(a)** Estimates of CH₄ enhancements from tower 8. **(b)** Estimates
 1242 of CH₄ enhancements from tower 13. **(c)** Estimates of CH₄ flux from tower 8. **(d)** Estimates of CH₄ flux
 1243 from tower 13.



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Figure 8. Google Earth image overlaid with bivariate polar plots (section 2.5) of the CH₄ enhancements at 9 INFLUX towers in Indianapolis using the criterion 1 background (Table 1) for full years of 2014 and 2015 over the afternoon (12-16 LST). The wind speed scale is only labeled at site 1; other sites follow the same convention. Legend indicates known sources of CH₄: Panhandle Eastern Pipeline (PEP), Southern Side Landfill (SSLF), Twin Bridges Landfill (TBLF), Waste Management Solutions (WMS), and Waste Water treatment facility (WW). The known magnitudes of sources that are in Marion County (PEP, SSLF, and WW) are reported in section 2.7. Magnitudes of TBLF and WMS according to EPA are approximately 5 mol/s. The largest known source on the map is SSLF.



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Figure 9. Google Earth image overlaid with bivariate polar plots (section 2.5) of the CH₄ enhancements at 9 INFLUX towers in Indianapolis using the criterion 1 background (Table 1) for year 2016 over the afternoon (12-16 LST). The wind speed scale is only labeled at site 1; other sites follow the same convention. Legend indicates known sources of CH₄: Panhandle Eastern Pipeline (PEP), Southern Side Landfill (SSLF), Twin Bridges Landfill (TBLF), Waste Management Solutions (WMS), and Waste Water treatment facility (WW). The known magnitudes of sources that are in Marion County (PEP, SSLF, and WW) are reported in section 2.7. Magnitudes of TBLF and WMS according to EPA are approximately 5 mol/s. The largest known source on the map is SSLF.