

1 **Response to Editor**

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We thank the editor for the robust examination of our paper and useful comments.

Comments to the Author:

Thank you for addressing the issues raised.

I have some minor comments below, all about wording.

46 ...estimate the magnitude...

53 you might want to add a reference to the recent paper by Hmiel et al (https://doi.org/10.1038/s41586-020-1991-8).

149 ...that, based on the EPA gridded inventory (Maasackers et al, 2016), increase...

170 Wind speed and direction were measured...

273ff Temporal variability in what? Background concentration, emission?? Do not write much but do clarify.

337 Inventory

422 ...using the INFLUX network, we estimate that... (2 changes)

444 ...features in Figure 7 is a ...

512methods. The atmospheric inverse flux..... (Could be Our atmospheric....)

All of the comments above were addressed to the best of our ability (see the updated combined document).

37 **Background Heterogeneity and Other Uncertainties in**
38 **Estimating Urban Methane Flux: Results from the**
39 **Indianapolis Flux (INFLUX) Experiment**
40

41 Nikolay V. Balashov^{1*,2,3}, Kenneth J. Davis¹, Natasha L. Miles¹, Thomas
42 Lauvaux^{1,4}, Scott J. Richardson¹, Zachary R. Barkley¹, Timothy A. Bonin^{5,6}
43

44 ¹The Pennsylvania State University, University Park, Pennsylvania, USA

45 ²NASA Postdoctoral Program, Universities Space Research Association, 7178 Columbia Gateway Drive,
46 Columbia, MD, 21046, USA

47 ³NASA Global Modeling and Assimilation Office (GMAO), Goddard Space Flight Center, Greenbelt,
48 MD, 20771, USA

49 ⁴Laboratory of Climate Sciences and Environment, Gif-sur-Yvette, France

50 ⁵Cooperative Institute for Research in Environmental Sciences, Boulder, Colorado, USA

51 ⁶Chemical Sciences Division, National Oceanic and Atmospheric Administration, Boulder, Colorado,
52 USA

53 *Former affiliation

54 *Correspondence to:* Nikolay V. Balashov (nvb5011@psu.edu or nikolay.v.balashov@nasa.gov) and Kenneth J.
55 Davis (kjd10@psu.edu)

56 **Abstract**

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

69 remains about 91% higher than inventory estimates. Hourly urban background CH₄ mole
70 fractions are shown to be spatially heterogeneous and temporally variable. Variability in
71 background mole fractions observed at any given moment and a single location could be up to
72 about 50 ppb depending on a wind direction, but decreases substantially when averaged over
73 multiple days. Statistically significant, long-term biases in background mole fractions of 2-5 ppb
74 are found from single point observations for most wind directions. Boundary layer budget
75 estimates suggest that Indianapolis CH₄ emissions did not change significantly when comparing
76 2014 to 2016. However, it appears that CH₄ emissions may follow a diurnal cycle with daytime
77 emissions (12-16 LST) approximately twice as large as nighttime emissions (20-5 LST). We
78 found no evidence for large CH₄ point sources that are otherwise missing from the inventories.
79 The data from the towers confirm that the strongest CH₄ source in Indianapolis is South Side
80 Landfill. Leaks from the natural gas distribution system that were detected with the tower
81 network appeared localized and non-permanent. Our simple atmospheric budget analyses
82 estimate [the](#) magnitude of the diffuse NG source to be 70% higher than inventory estimates, but
83 more comprehensive analyses are needed. Long-term averaging, spatially-extensive upwind
84 mole fraction observations, mesoscale atmospheric modeling of the regional emissions
85 environment, and careful treatment of the times of day are recommended for precise and accurate
86 quantification of urban CH₄ emissions.

87

88 **1 Introduction**

89 From the beginning of the Industrial Revolution to 2011, atmospheric methane (CH₄) mole
90 fractions increased by a factor of 2.5 due to anthropogenic processes such as fossil fuel
91 production, waste management, and agricultural activities (Ciais et al., 2013; [Hmiel et al., 2020](#)).

92 The increase in CH₄ is a concern as it is a potent greenhouse gas (GHG) with a global warming
93 potential 28-34 times greater than that of CO₂ over a period of 100 years (Myhre et al., 2013).
94 The magnitudes of component CH₄ sources, and the causes of variability in the global CH₄
95 budget are not well understood with theories attributing changes in biogenic, thermogenic, and
96 pyrogenic emissions or a decline in the atmospheric CH₄ sink to the recent CH₄ increases (Nisbet
97 et al., 2016; Saunio et al., 2016; Nisbet et al., 2019; Hmiel et al., 2020). Improved
98 understanding of CH₄ emissions is needed (National Academies of Sciences and Medicine,
99 2018).

100 In particular, the estimates of continental U.S. anthropogenic CH₄ emissions disagree.
101 Inventories from Environment Protection Agency (EPA) and Emissions Database for Global
102 Atmospheric Research (EDGAR) in 2008 reported emission values of 19.6 and 22.1 TgC y⁻¹
103 (U.S. EPA, 2013; European Commission Joint Research Centre and Netherlands Environmental
104 Assessment Agency, 2010). However, top-down methodologies using aircraft and inverse
105 modeling framework found emission values of 32.4 ± 4.5 TgC y⁻¹ for 2004 and 33.4 ± 1.4 TgC
106 y⁻¹ for 2007-2008 respectively (Kort et al., 2008; Miller et al., 2013). Underestimation of natural
107 gas (NG) production and agricultural sources are possible reasons for this disagreement (Miller
108 et al., 2013; Brandt et al., 2014; Jeong et al., 2014). Efforts to reconcile GHGs emissions
109 estimates using atmospheric methods and inventory assessment have sometimes succeeded
110 (Schuh et al., 2013; Zavala-Araiza et al., 2015; Turnbull et al., 2019) when careful attention is
111 given to the details of each method, and targeted atmospheric data are available. A recent
112 synthesis of emissions from the U.S. NG supply chain demonstrated similar success and
113 concluded that current inventory estimates of emissions from U.S. NG production are too low

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118 and that emission from NG distribution is one of the greatest remaining sources of uncertainty in
119 the NG supply chain (Alvarez et al., 2018).

120 Due to the uncertainties in CH₄ emissions from NG distribution it is natural that urban
121 emissions are of interest as well. For example, two studies (McKain et al., 2015; Hendrick et al.,
122 2016) indicate that ~60-100% of Boston CH₄ emissions are attributable to the NG distribution
123 system. Recent studies of urban CH₄ emissions in California indicate that the California Air
124 Resources Board (CARB) inventory tends to underestimate the actual CH₄ urban fluxes possibly
125 due to fugitive emissions from NG infrastructures in urban environments (Wunch et al., 2009;
126 Jeong et al., 2016; Jeong et al., 2017). The accuracy and precision of atmospheric estimates of
127 urban CH₄ emissions are limited by available atmospheric observations (Townsend-Small et al.,
128 2012), potential source magnitude variability with time (Jackson et al., 2014; Lamb et al., 2016),
129 errors in atmospheric transport modeling (Hendrick et al., 2016; Deng et al., 2017; Sarmiento et
130 al., 2017), and complexity in atmospheric background conditions (Cambaliza et al., 2014; Karion
131 et al., 2015; Heimbürger et al., 2017). In this work, detailed analysis of urban CH₄ mole
132 fractions is performed for the city of Indianapolis to better understand the aforementioned
133 uncertainties of urban CH₄ emissions.

134 The Indianapolis Flux Experiment (INFLUX; Davis et al., 2017) is a testbed for
135 improving quantification of urban GHGs emissions and their variability in space and time.
136 INFLUX (<http://influx.psu.edu>) is located in Indianapolis partly because of its isolation from
137 other urban centers and the flat Midwestern terrain. It includes a very dense GHGs monitoring
138 network, comprised of irregular insitu aircraft measurements (Heimbürger et al., 2017;
139 Cambaliza et al., 2014), continuous in situ observations from communications towers using
140 cavity ring-down spectroscopy (Richardson et al., 2017; Miles et al., 2017), and automated flask

141 sampling systems for quantification of a wide variety of trace gases (Turnbull et al., 2015).
142 Meteorological sensors include a Doppler lidar providing continuous boundary layer depth and
143 wind profiles, and tower-based eddy covariance measurements of the fluxes of momentum,
144 sensible and latent heat (Sarmiento et al., 2017). The network is designed for emissions
145 quantification using top-down methods such as tower-based inverse modeling (Lauvaux et al.,
146 2016) and aircraft mass balance estimates (Cambaliza et al., 2015).

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

156 Biogenic emissions from the city are dominated by a landfill close to downtown, and
157 these emissions are thought to be fairly well known (GHG reporting program). Although
158 evidence of possible variability in landfill emissions exists from Cambaliza et al. (2015), which
159 used aircraft mass balance on five different occasions to calculate CH₄ flux from this landfill.
160 Uncertainty in total city emissions is mainly driven by the uncertainty in thermogenic emissions,
161 which are hypothesized to emerge largely from the NG distribution system (Mays et al., 2009;
162 Cambaliza et al., 2015; Lamb et al., 2016). In this study, we explore potential explanations for

163 the discrepancies in CH₄ emissions estimates from Indianapolis and posit methods and
164 recommendations for the study of CH₄ emissions from other urban centers.

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

172

173 **2 Methods**

174

175 **2.1 Experimental site**

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

184 Figure 2 shows two domains that have been used for the evaluation of Indianapolis CH₄
185 emissions (Lamb et al., 2016; Lauvaux et al., 2016). The first domain is the whole area shown in

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187 the figure enclosing both Indianapolis and places that lie outside of its boundaries. This domain
188 was used for the inversion performed in Lamb et al. (2016). The second domain is Marion
189 County outlined with a green dashed line. It is assumed here that this domain is much more
190 representative of the actual Indianapolis municipal boundaries as this area encompasses the
191 majority of the urban development associated with the city of Indianapolis (Gurney et al., 2012).
192 The larger domain has three additional landfills that, based on the EPA gridded inventory
193 (Maasakkers et al., 2016), increase Indianapolis CH₄ emissions by about 50% when compared to
194 the smaller domain. The inversion explained in Lamb et al. (2016) has been rerun for two of the
195 domains mentioned above and the results (Fig. 1) have been reexamined.

196

197 **2.2 INFLUX tower network**

198 The continuous GHG measurements from INFLUX are described in detail in Richardson et al.
199 (2017). The measurements were made using wavelength-scanned cavity ring down
200 spectrometers (CRDS, Picarro, Inc., models G2301, G2302, G2401, and G1301), installed at the
201 base of existing communications towers, with sampling tubes secured as high as possible on each
202 tower (39 – 136 m above ground level (AGL); Miles et al., 2017). A few towers also included
203 measurements at 10 m AGL and one or two intermediate levels. While INFLUX tower in-situ
204 measurements began in September 2010, here we focus on the CH₄ measurements from 2013 –
205 2016. From June through December 2012, there were two or three towers with operational CH₄
206 measurements. By July 2013, five towers included measurements of CH₄, and throughout the
207 majority of the years 2015 – 2016 there were eight INFLUX towers with CH₄ measurements
208 (Fig. 3). Flask to in-situ comparisons and round-robin style testing indicated compatibility

209 across the tower network of 0.6 ppb CH₄ (Richardson et al., 2017). In this study we use hourly
210 means of CH₄.

211

212 2.3 Meteorological data

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

223 The planetary boundary layer height (BLH) was determined from a Doppler lidar
224 deployed in Lawrence, Indiana about 15 km to the northeast of downtown. The lidar is a Halo
225 Streamline unit, which was upgraded to have extended range capabilities in January 2016. The
226 lidar continuously performs a sequence of conical, vertical-slice, and staring scans to measure
227 profiles of the mean wind, turbulence, and relative aerosol backscatter. All of these
228 measurements are combined using a fuzzy-logic technique to automatically determine the BLH
229 continuously every 20-min (Bonin et al., 2018). The BLH is primarily determined from the
230 turbulence measurements, but the wind and aerosol profiles are also used to refine the BLH
231 estimate. The BLHs are assigned quality-control flags that can be used to identify times when

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233 the determined BLH is unreliable, such as when the air is exceptionally clean, the BLH is below
234 a minimum detectable height, or clouds and fog that attenuate the lidar signal exist. Additional
235 details about the algorithm and the lidar operation for the INFLUX project are provided in Bonin
236 et al. (2018). Doppler lidar measurements are available at
237 <https://www.esrl.noaa.gov/csd/projects/influx/>.

238

239 **2.4 Urban methane background**

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

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

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

250 Aircraft mass balance studies of Indianapolis mentioned used two main methods to
251 determine a background value. The first method calculates an average of the aircraft transect
252 edges that lie outside of the city domain (Cambaliza et al., 2014). In the second approach, a
253 horizontally varying background is introduced by linearly interpolating median background
254 values of each of the transect edges (Heimbürger et al., 2017). In theory there is also a third

255 method that uses an upwind transect as a background field, but in the studies above it was
256 assumed that the edges are representative of an upwind flow. In the case of an inversion, it is
257 common to pick a tower that is located generally away from urban sources and has on average
258 the smallest overall enhancement (Lavaux et al., 2016). Because choosing the background
259 involves a degree of subjectivity (Heimburger et al., 2017) we consider how these choices may
260 influence emission estimates and introduce error, both random and systematic, using data from
261 the INFLUX tower network.

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

272 Criterion 1 corresponds to a typical choice of a background in a case of tower inversion
273 and is based on the concept that the lowest CH₄ mole fraction measured at any given time is not
274 affected by the city sources and therefore is a viable approximation of the background CH₄ mole
275 fractions outside of the city (Miles et al., 2017; Lauvaux et al., 2016). Given this assumption, the
276 tower with the lowest median of the CH₄ enhancement distribution (calculated by assuming the
277 lowest measurement among all towers at a given hour as a background) for each of the wind

278 directions over the November 2014 through December 2016 time period is chosen as a
279 background site (Miles et al., 2017). Criterion 2 requires that the tower is outside of Marion
280 County (outside of the city boundaries) and is not downwind of any known regional CH₄ source
281 (Fig. 2). For some wind directions, there are multiple towers that could qualify as a background;
282 we pick towers in such a manner that they are different for each criterion given a wind direction
283 in order to calculate the error associated with the use of different but acceptable backgrounds.
284 The towers used for both criteria and for each of the eight wind directions are displayed in Table
285 1. Quantifying differences between these two backgrounds allows for an opportunity to better
286 understand the degree of uncertainty that exists in the atmospheric CH₄ background at
287 Indianapolis.

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

299

300 **2.5 Frequency and bivariate polar plots**

301 Frequency and bivariate polar plots are used in this work to gain more knowledge regarding CH₄
302 background variability based on criteria 1 and 2, and to identify sources located within the city.
303 To generate these polar plots, we use the *openair* package (from R programming language)
304 created specifically for air quality data analysis (Carslaw and Ropkins, 2012). Bivariate and
305 frequency polar plots indicate the variability of a pollutant concentration at a receptor (such as an
306 observational tower) as a function of wind speed and wind direction, preferably measured at the
307 location of the receptor or within several kilometers of the receptor. The frequency polar plot is
308 generated by partitioning the CH₄ hourly data into the wind speed and direction bins of 1 m s⁻¹
309 and 10° respectively. To generate bivariate polar plots, wind components *u* and *v* are calculated
310 for hourly CH₄ mole fraction values, which are fitted to a surface using a Generalized Additive
311 Model (GAM) framework in the following way,

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

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

316

317 2.6 Temporal variability and approximate flux estimation

318 Temporal variability in urban CH₄ emissions may play an important role in the corresponding
319 emissions quantification procedures. Lamb et al. (2016) suggested that such temporal variability
320 might partially explain the differences among CH₄ flux estimates shown in Figure 1. If temporal
321 variability of CH₄ emissions exists within the city, disagreements in the CH₄ flux between
322 studies could be attributed to differences in their sampling period. Because the INFLUX tower

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325 data at Indianapolis contain measurements at all hours of the day over multiple years, we can
326 utilize this dataset to better understand the temporal variability in methane emissions in the city.

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

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

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

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

342 We use equation 4 to estimate hourly CH₄ emissions (Q_a) from Indianapolis (see
343 assumptions in the paragraph below) given hourly averaged data of H from the lidar positioned
344 in the city, wind speed (u) from the local weather stations, and upwind (C_b) and downwind (C)

345 CH₄ mole fractions measured (and then converted to concentrations) at towers 1, 8, and 13
346 (depending on a wind direction) using data from heights of 40 m, 41 m, and 87 m respectively
347 (see Fig. 2).

348 The CH₄ concentrations are derived from CH₄ mole fractions by approximating average
349 molar density of dry air (in mol m⁻³) within the boundary layer for every hour of the day, where
350 variability of pressure with altitude is calculated using barometric formula and it is assumed that
351 temperature decreases with altitude by 6.5 K per kilometer. The hourly surface data for pressure
352 and temperature are taken from KIND weather station. The difference between concentrations
353 $C - C_b$ is instantaneous and not lagged, where C_b represents air parcel entering the city and C
354 represents the same air parcel exiting the city (Turnbull et al., 2015). The CH₄ enhancements
355 $C - C_b$ are estimated for daytime by averaging observations spanning 12-16 LST and for
356 nighttime by averaging observations spanning 20-5 LST. These time periods are based on lidar
357 estimations of when on average H varies the least. The day and night were required to contain at
358 least 3 and 9 hourly CH₄ values respectively for averaging to occur, otherwise the day/night is
359 eliminated. Observations when H is below 100 m are not used to avoid the cases when
360 measurements from towers may be above the boundary layer. In order to better achieve the
361 assumption that the boundary layer is fully mixed (especially at night), all hours with wind
362 speeds below 4 m/s are eliminated (Van De Wiel., 2012). To approximate the emissions of the
363 whole city we need to know the approximate area of the city and the distance over which the
364 plume is affected by the city CH₄ sources. The area of the city is about 1024 km² (the area of
365 Marion County) and the length that plume traverses when it is over the city ranges from 32 to 35
366 km depending on which downwind tower is used. We assume that CH₄ measurements at towers
367 8 and 13 are representative of a vertically well-mixed city plume as the towers are located

368 outside of the city boundaries and allow for sufficient vertical mixing to occur. For S and SW
369 wind directions tower 8 observations are used to represent downwind conditions with
370 background observations coming from towers 1 and 13, respectively (based on criterion 1 shown
371 in Table 1). For W wind direction, tower 13 observations represent the downwind with
372 background obtained from tower 1. The wind direction is required to be sustained for at least 2
373 hours, otherwise the data point is eliminated.

374

375 2.7 Indianapolis CH₄ sources

376 Only a few known CH₄ point sources exist within Indianapolis (Cambaliza et al., 2015; Lamb et
377 al., 2016). The Southside Landfill (SSLF), located near the center of the city, is thought to be the
378 largest point source in the city with emissions ranging between about 28 mol/s (inventory from
379 Maasackers et al. (2016), GHG reporting program, and inverse estimates from ground-based
380 mobile sampling employed in Lamb et al. (2016)) and 45 mol/s (aircraft; Cambaliza et al.
381 (2015)) depending on an emission estimation methodology. However, using Cambaliza et al.
382 (2015) aircraft data and applying a different background formulation Lamb et al. (2016) found
383 emission values of SSLF closely agreeing with 28 mol/s estimate. SSLF could account for as
384 little as 33% (top-down from Cambaliza et al. (2015)) or as much as 63% (inventory from
385 Maasackers et al. (2016)) of total Marion County CH₄ emissions. Other city point sources are
386 comparatively small; the wastewater treatment facility located near SSLF contributes about 3-7
387 mol/s (inventory from Lamb et al. (2016)), and the transmission-distribution transfer station at
388 Panhandle Eastern Pipeline (also known as a city gate and further in this study abbreviated as
389 PEP) is estimated to be about 1 mol/s (inventory from Lamb et al. (2016)). The remaining CH₄
390 sources, mainly from NG infrastructure leaks and livestock, are considered to be diffuse sources

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395 and are not well known. Potential sources of emissions related to NG activities include gas
396 regulation meters, transmission and storage, distribution leaks, and Compressed Natural Gas
397 (CNG) fleets. These diffuse NG sources account for 21-67% of the city emissions or 20 mol/s
398 (inventory from Maasackers et al., (2016)) to 64 mol/s (top down from Cambaliza et al., (2015)).
399 Livestock emissions for Marion County are estimated to be around 1.5 mol/s (inventory from
400 Maasackers et al., (2016)). These prior studies present conflicting conclusions regarding the
401 magnitude of the diffuse NG CH₄ source in Indianapolis.

402

403 3 Results and discussion

404

405 3.1 Inversion and city boundaries

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

412 In the case of Indianapolis, this issue became apparent when the emissions were
413 calculated using an atmospheric inversion model (Lamb et al., 2016; Lauvaux et al., 2016). The
414 atmospheric inversion solved for fluxes in domain 1 (Fig. 2), which significantly increased the
415 estimated emissions in comparison with the inventory values that were gathered mainly within
416 Marion County (domain 2). When reduced to domain 2, inverse modeling emission estimate
417 decreases to 107 mol/s (from about 160 mol/s), which falls within an error bar of Lamb et al.
418 (2016) inventory estimate. This difference is significant and could at least partially explain the
419 discrepancy shown in Figure 1 between the emission values from the inventories and emission

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

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

429

430 **3.2 Variability in CH₄ background**

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

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

446 criterion 1 from November 2014 to December 2016 for each of the eight wind directions
447 mentioned in Table 1. The results are shown in Figure 5. Systematic bias is evident for the SE,
448 S, SW, W, and NW wind sectors, whereas random error dominates N, NE, and E wind
449 directions. Wind directions showing statistically significant bias have mean biases ranging from
450 2 to 5 ppb, with values as large as 8 ppb falling within the range of $2 \times$ standard error. Standard
451 deviation plot indicates potential background discrepancy that can occur on any given day, where
452 W wind direction is the least variable with $2 \times$ standard deviation close to 20 ppb, while SE wind
453 direction is the most variable with $2 \times$ standard deviation falling at about 50 ppb.

454 Random errors in the mole fractions of background differences (biases) are also
455 important and are a function of the length of the data record. We quantify the random error in
456 the CH₄ background mole fraction differences using the bootstrap method by randomly sampling
457 2 to 150 hours (small and large sample size) of the background CH₄ differences for each of the
458 wind directions with replacement (we make the assumption that our differences are independent
459 since we eliminated lag 1 autocorrelation from the data). This sub-sampling experiment is
460 repeated 5000 times (Efron and Tibshirani, 1986). The standard deviations of the mean
461 (standard error) of the 5000 simulated differences are calculated for each wind direction. The
462 resulting standard errors of the city CH₄ background differences, multiplied by 2 to represent the
463 95% confidence intervals, are shown as a function of the length of the data record in Figure 6.
464 Because random error falls as sample size grows it makes sense to assign a threshold indicating a
465 minimum number of samples needed to achieve a theoretical precision for each wind direction.

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

469 combined with a bias (Table 2). Meaning that the sum of bias and standard error must be at most
470 6 ppb. In this approach each wind direction would have a different threshold because of the
471 differences in biases. For instance, given this requirement NW direction would need a random
472 error of 1 since its bias is 5. For NW direction, this threshold would require more than 150
473 samples. For N direction on the other hand, where the bias is 1, the requirement is fulfilled when
474 random error crosses 5 ppb at 74 samples. Now we consider these random and systematic errors
475 in CH₄ background differences in the context of Indianapolis urban CH₄ emissions.

476 | For Indianapolis, using [the INFLUX network](#), we estimated that depending on sample
477 size (number of hours sampled) and wind direction, background gradient across the city over 12-
478 16 LST could vary from 0 to about 50 ppb (Fig. 5b). Given that the average afternoon CH₄
479 enhancement of the city is around 8-12 ppb (section 3.3; Fig. 7; Cambaliza et al., 2015; Miles et
480 al., 2017), the error on the estimated emissions could easily be over 100% if the analysis does not
481 approach the issue of background with enough sampling. A sample size of about 50 independent
482 hours significantly decreases background uncertainty for N, NE, E, S, and W wind directions and
483 allows for a more accurate assessment of the CH₄ emissions at Indianapolis. For CH₄ sources
484 with a significantly larger signal than their regional background, the mentioned background
485 variability becomes less impactful on results, but because Indianapolis is a relatively small
486 emitter of CH₄, and because there are relatively large sources outside of the city, uncertainties
487 due to background estimation are comparatively large. Our uncertainty assessment suggests that
488 the highly variable CH₄ emission values of Indianapolis from aircraft mass balance calculations
489 shown in Figure 1 are at least partially due to the variability in the urban CH₄ background of
490 Indianapolis.

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493 3.3. Temporal variability of methane enhancements and fluxes in Indianapolis

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

499 One of the more interesting features in Figure 7 is a day/night variability of CH₄
500 emissions at Indianapolis. The most prominent example of this feature is found in Figure 7c,
501 where the estimates for both years suggest that daytime emissions are approximately twice as
502 large as the emissions at night. The decrease of the CH₄ emissions at night also appears in tower
503 13, but the errors are too high in those estimates to make any definitive conclusions. A similar
504 urban CH₄ emissions diurnal variability is reported by Helfter et al. (2016) in their study of
505 GHGs for London, UK, where they attribute diurnal variation of CH₄ emissions to the NG
506 distribution network activities, fugitive emissions from NG appliances, and to temperature-
507 sensitive CH₄ emission sources of biogenic origin (such as a landfill). Taylor et al. (2018)
508 suggest that CH₄ emissions from landfills exhibit a diurnal cycle with higher emissions in early
509 afternoon and 30-40% lower emissions at night.

510 With regard to yearly temporal variability we are only able to compare years 2014 and
511 2016 due to limited BLH data for other years. Results from both towers suggest that
512 Indianapolis overall CH₄ emissions did not change significantly between 2014 and 2016.
513 Although it is important to be cautious about interpreting actual flux estimations given the
514 assumptions mentioned in section 2.6, it is interesting to note that the flux values from both
515 towers average to about 70 mol/s, which puts our value right in between inventory and inversion

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517 estimates shown in Figure 1. If we assume that SSLF emissions are generally known (GHG
518 reporting program) that would indicate that emissions from NG distribution are likely to be about
519 14 mol/s (70%) higher than what both of the inventories currently estimate but within the error
520 bars of Lamb et al. (2016)'s inventory calculation. Another possible scenario is that SSLF
521 emissions are higher than what is currently assumed. Given these complexities, uncertainty
522 regarding the exact emissions from NG distribution at Indianapolis still remains.

523

524 **3.4 Methane Sources in Indianapolis**

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

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

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

548 Another important point is the cluster of large enhancements surrounding tower 10 in
549 2014-2015. Because no other tower sees these enhancements (at least at comparable
550 magnitudes), we believe that they are the result of nearby NG leaks. These plumes are not
551 consistent temporally or spatially as they mostly disappear in 2016, potentially indicating that
552 they are transient and localized NG distribution leaks. It is difficult to ascertain the exact
553 combined magnitude of these leaks since they mix together with SSLF into an aggregated city
554 plume when observed from downwind towers such as 8 and 13. None of the individual leaks
555 appears to be similar in magnitude to the emissions that originate from SSLF. Diffuse NG
556 emissions comparable to the SSLF source (Lamb et al., 2016) may exist. Our flux estimations at
557 towers 8 and 13, however, imply that the magnitude of NG diffuse source suggested by the top-
558 down analyses in Cambaliza et al. (2015) and Lamb et al. (2016) are probably overestimates (see
559 section 3.3). We hypothesize that the relatively high Indianapolis CH₄ emissions (see Fig. 1)
560 reported by Cambaliza et al. (2015) could be a result of random errors in upwind conditions (see
561 section 3.2) influencing the small number of airborne flux estimates.

562

563 4 Conclusions

564 We have examined four potential contributions to discrepancies between urban top-down and
565 bottom-up estimates of CH₄ emissions from Indianapolis: domain definition, heterogeneous
566 background mole fractions, temporal variability in emissions, and sources missing from
567 inventories. Results indicate that the urban domain definition is crucial for the comparison of the
568 emission estimates among various methods. Our atmospheric inverse flux estimates for Marion
569 County, which is similar to the domain that is analyzed by inventory and airborne mass balance
570 methodologies (Mays et al., 2009; Cambaliza et al., 2014; Lamb et al., 2016), is 107 mol/s
571 compared to 160 mol/s that is estimated for the larger domain (Hestia inventory domain; Gurney
572 et al., 2012). This partially explains higher emissions in inverse modeling estimates shown by
573 Lamb et al. (2016); however, 107 mol/s is still 91% higher than what EPA and Lamb et al.
574 (2016) find in their inventories (Fig. 1).

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

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589 to random errors on the order of 10-30 ppb in the CH₄ background, similar to the magnitude of
590 the total enhancement from the city of Indianapolis, which is estimated to be on average around
591 10-12 ppb. Longer-term sampling and/or more extensive background sampling are necessary to
592 reduce the random errors. Sample size required to reduce random errors of background
593 differences to an acceptable value for flux calculation is largely dependent on a wind direction.
594 Both bias (long-term average of background differences) and its random error are important
595 when estimating total background uncertainty. The results indicate that N, NE, E, S, and W
596 wind directions are more favorable for flux estimation and would require multiple days of
597 measurements (e.g. about 50 independent hours of measurements) to reduce background
598 uncertainty to about 6 ppb, which is half the magnitude of the typical CH₄ enhancement from
599 Indianapolis. The remaining wind directions would require over 150 independent hourly
600 measurements to achieve similar precision. We also estimate that depending on a wind direction
601 for any given hour the spatial variability in background can be anywhere from 0 to 50 ppb. This
602 uncertainty in the CH₄ background may partially explain Heimburger et al. (2017) finding of
603 large variability in airborne estimates of Indianapolis CH₄ emissions. Given many samples, the
604 airborne studies converge to an average value of CH₄ flux that is noticeably closer to the
605 inventory estimates for Indianapolis than several of the individual estimates presented in Figure
606 1.

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

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

618 With regard to temporal variability, no statistically detectable changes in the emission
619 rates were observed when comparing 2014 and 2016 CH₄ emissions. However, a large
620 difference between day and night CH₄ emissions was implied from a simple budget estimate.
621 Night (20-5 LST) emissions may be 2 times lower than the emissions during the afternoon (12-
622 16 LST) hours. Because prior estimates of top-down citywide emissions are derived using
623 afternoon-only measurements, overall emissions of Indianapolis may be lower than these studies
624 suggest. This bias may be present in studies performed in other cities as well. Our study
625 suggests that day/night differences in CH₄ emissions must be understood if regional emission
626 estimates are to be calculated correctly. Long-term, tower-based observations are an effective
627 tool for understanding and quantifying multi-year variability in urban emissions.

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

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

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

644

645 **Author Contribution**

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

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

658

659 **Competing Interests**

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

661

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669 and noticeably improved clarity of our article.

670

671

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907 **Tables**

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Table 1. INFLUX towers used to estimate CH₄ background based on two different criteria. Numbers in bold indicate towers chosen to generate a background field when multiple options are possible (for more details see discussion). In short, criterion 1 uses towers with the lowest mean CH₄ for a specific wind direction, and criterion 2 uses towers outside of Marion County and not downwind of large sources (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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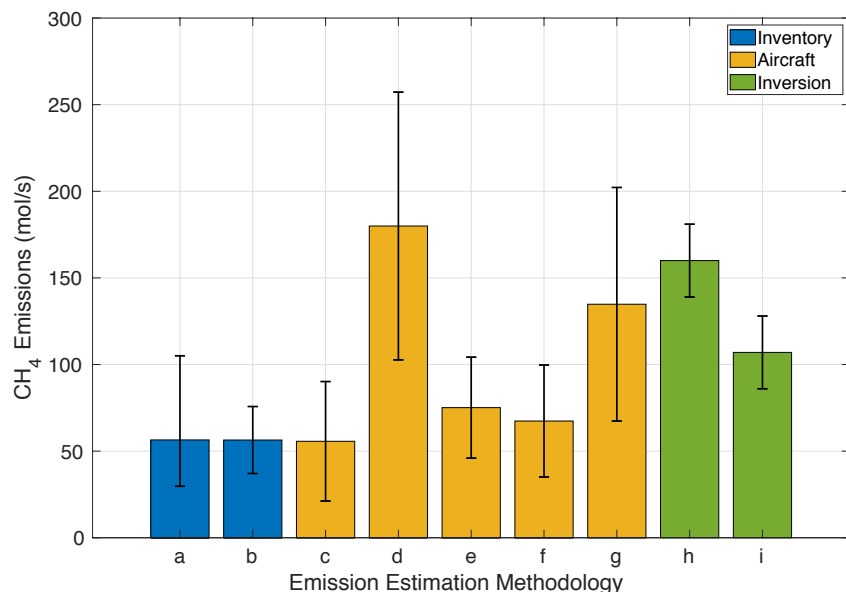
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931 **Table 2.** A number of independent samples needed (column 4) to satisfy combined requirement of 6 ppb
932 background error based on the sum of bias and random error (explained in section 3.2) as a function of
933 wind direction.

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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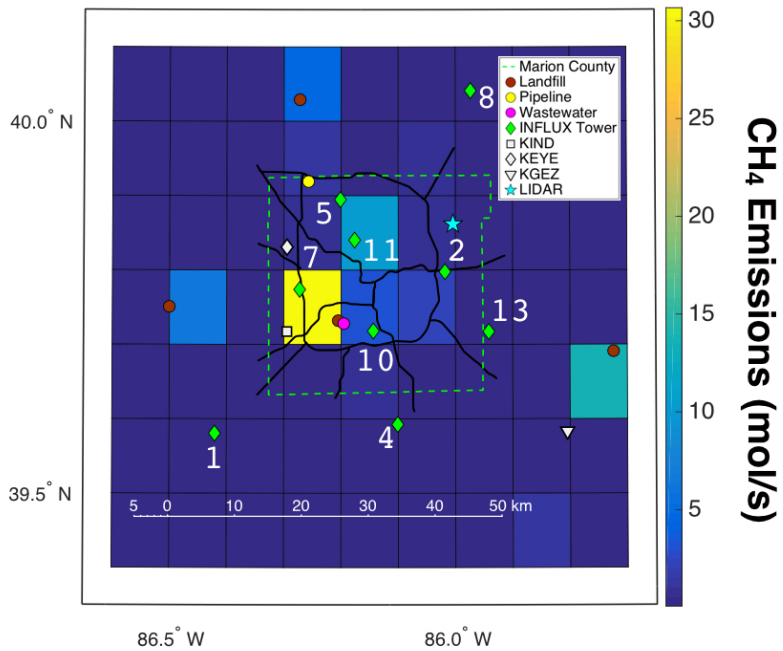
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962 **Figures**
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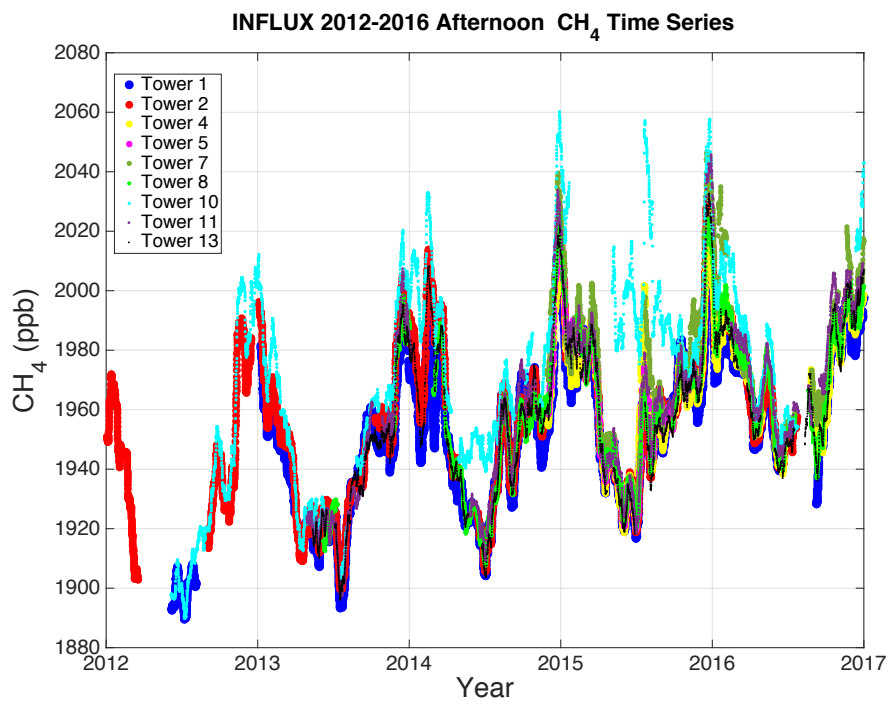
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965 **Figure 1.** Various estimates of CH₄ emissions at Indianapolis. **(a, b)** Bottom-up estimates of CH₄
966 emissions conducted by Lamb et al. (2016) in 2013 and Maasakkers et al. (2016) based on the EPA 2012
967 inventory respectively. Error bars show 95% confidence intervals (for more details see above-mentioned
968 articles). **(c-g)** Top-down evaluations of CH₄ emissions with aircraft from various flight campaigns where
969 **(c)** contains 5 flights over March-April of 2008, **(d)** contains 3 flights over November-January of 2008-
970 09, **(e)** contains 5 flights over April-July of 2011, **(f)** contains 9 flights from November-December, 2014,
971 and **(g)** contains the same 5 flights over April-July of 2011 as in (e) but uses different methodology.
972 Methodologies for **(c-f)** are described in Lamb et al. (2016) and methodology for **(g)** is described in
973 Cambaliza et al. (2015). Error bars show 95% confidence intervals (for more details see above-
974 mentioned articles). **(h, i)** Top-down evaluations of CH₄ emissions for 2012-2013 using tower inversion
975 modeling methodology with two different domains, where **(h)** uses the full domain of Figure 2 and **(i)**
976 uses only the Marion County domain of Figure 2. The inversion methodology and 95% confidence
977 intervals are described in detail in Lamb et al. (2016).

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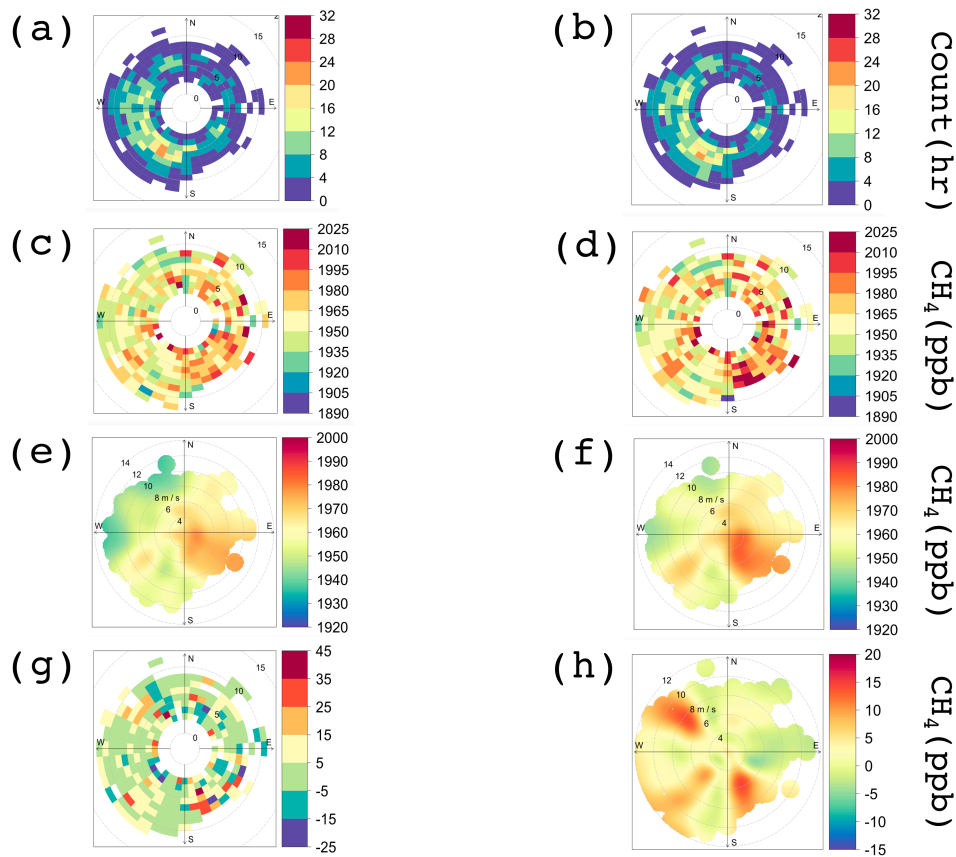


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 983 **Figure 2.** Map of the primary roads in Indianapolis, INFLUX towers, lidar system, weather stations, and
 984 a few CH₄ point sources plotted over the gridded CH₄ emissions (mol/s) from the EPA 2012 Inventory
 985 (Maasackers et al., 2016). The gridded map of emissions includes emissions from the mentioned point
 986 sources; their position is provided to aid in interpretation of the observations. The dashed bright green
 987 line denotes Marion County borders.
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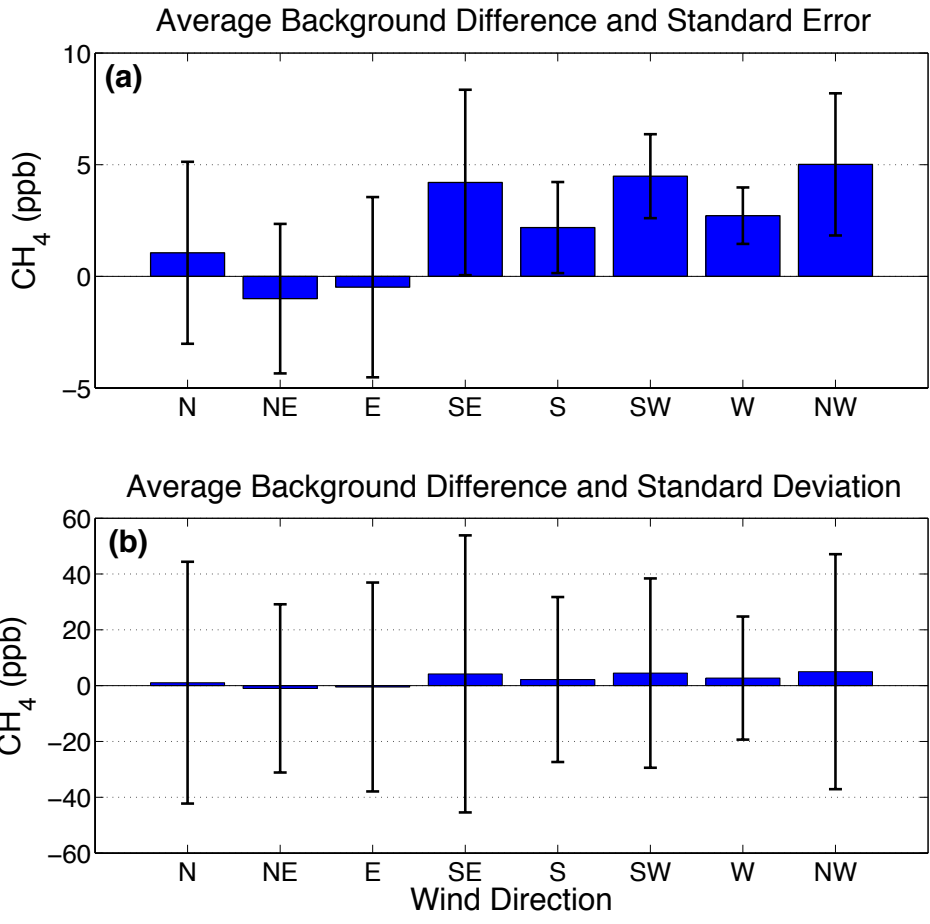
990 **Figure 3.** 20-day running average of afternoon (12-16 LST; the hours are inclusive) CH₄ mole fractions
 991 as measured by the INFLUX tower network (highest available height is used) from 2012 through 2016.



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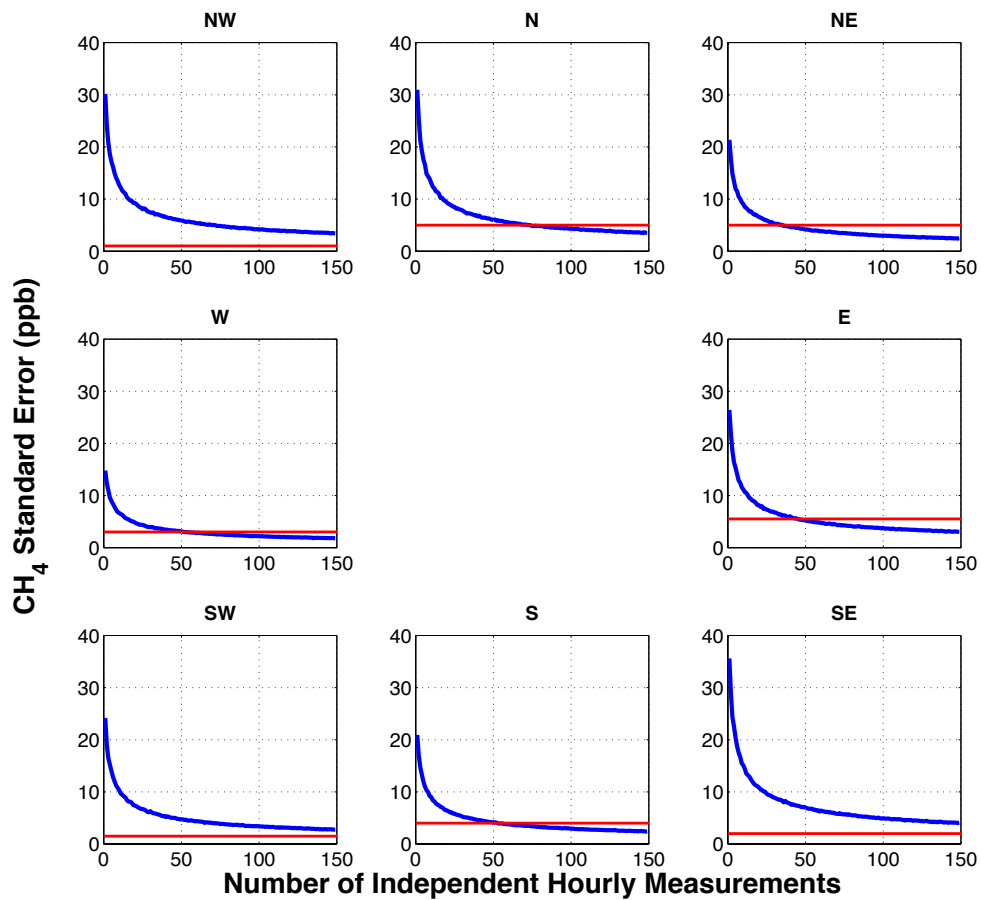
993 **Figure 4.** Frequency and bivariate polar plots of CH₄ background for Indianapolis using data from 12-16
 994 LST, November 2014 through December 2016 given 2 different criteria (Table 1). **(a)** Polar histogram
 995 indicating a number of hourly measurements available using criterion 1. **(b)** Same as (a) only for criterion
 996 2. Differences between (a) and (b) are due to slight differences in data availability at the considered
 997 towers. **(c)** Polar frequency plot of the CH₄ background using criterion 1. **(d)** Same as (c) only for
 998 criterion 2. **(e)** Polar bivariate plot of CH₄ background using criterion 1. **(f)** Same as (e) only for criterion
 999 2. **(g)** Polar frequency plot of difference between the backgrounds: *criterion 2 – criterion 1*. **(h)** Same
 1000 as (g) but shown with a bivariate polar plot.

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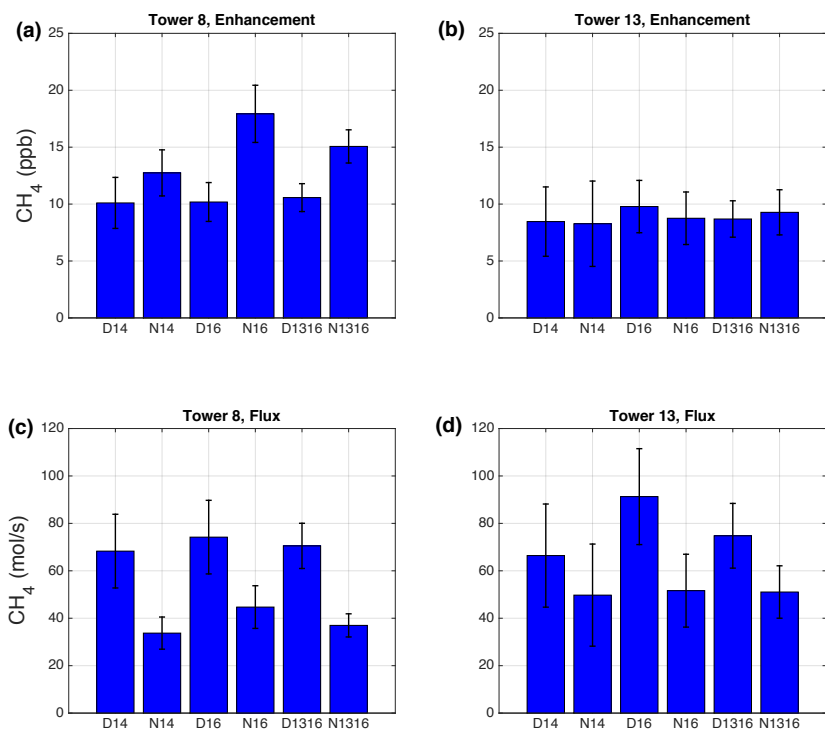
1002 **Figure 5.** Average of the differences between criteria 2 and 1 CH₄ backgrounds at Indianapolis as a
 1003 function of wind direction. These averages are generated from the same data that is used in Figure 4 and
 1004 reflect results shown in Figure 4g. Error bars indicate in (a) 2 × standard error and in (b) 2 × standard
 1005 deviation.
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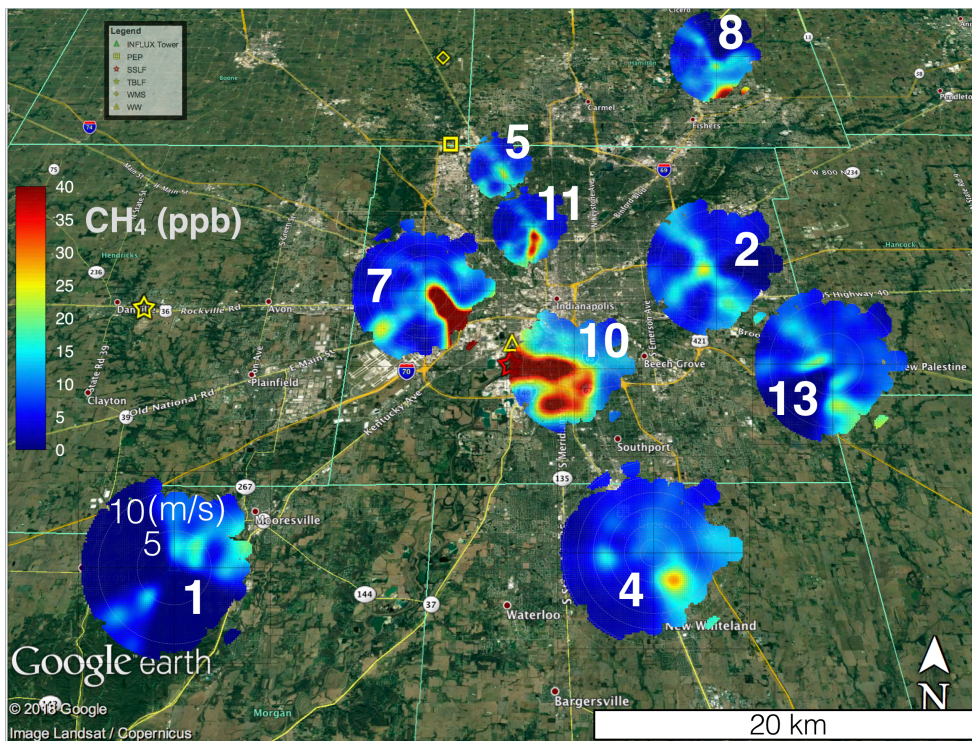
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Figure 6. Bootstrap simulation of the standard errors $\times 2$ in Indianapolis CH₄ background mole fraction differences (between criteria 2 and 1) as a function of sample size and wind direction (see text for details). Thresholds for each of the wind directions indicate a random error threshold needed for the background uncertainty to be within 50% of Indianapolis CH₄ enhancement of 12 ppb.



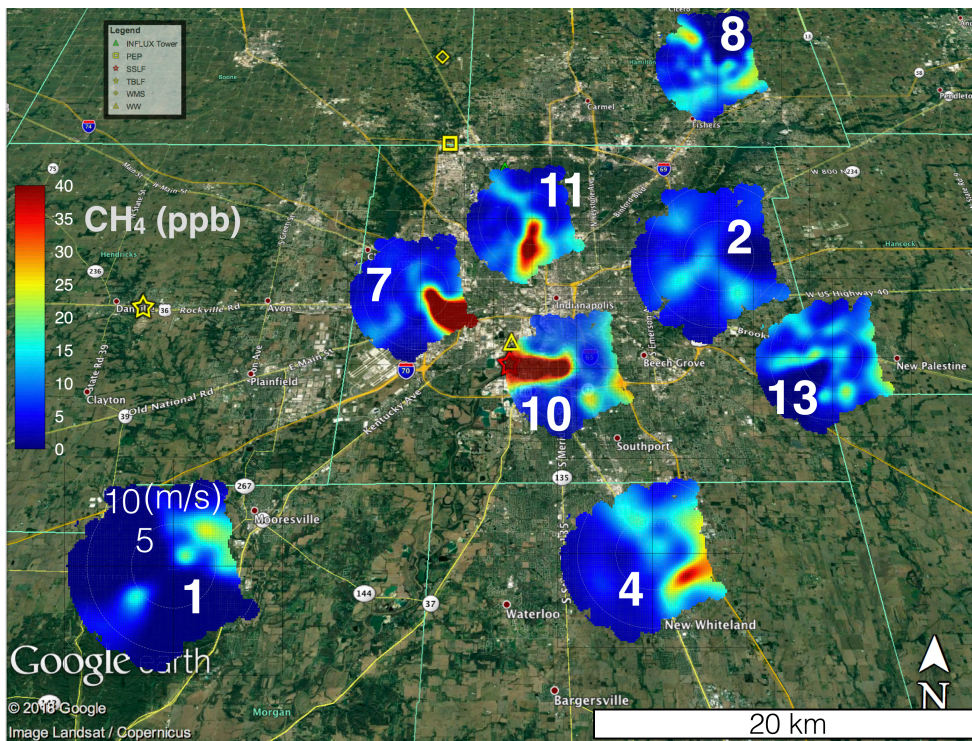
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1015 **Figure 7.** Averages of the daytime (D) and nighttime (N) CH₄ enhancements and fluxes at INFLUX
 1016 towers 8 and 13 for years 2014 (14), 2016 (16), and 2013-2016 (1316). The error bars represent 95%
 1017 confidence interval of each mean value. **(a)** Estimates of CH₄ enhancements from tower 8. **(b)** Estimates
 1018 of CH₄ enhancements from tower 13. **(c)** Estimates of CH₄ flux from tower 8. **(d)** Estimates of CH₄ flux
 1019 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.