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Daniel Zavala Araiza

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The Dissertation Committee for Daniel Zavala Araiza certifies that this is the approved version of the following dissertation:

**Atmospheric Emissions and Air Quality Impacts of Natural
Gas Production from Shale Formations**

Committee:

David Allen, Supervisor

Michael Webber

Elena McDonald-Buller

Lea Hildebrandt Ruiz

Thomas Edgar

**Atmospheric Emissions and Air Quality Impacts of Natural
Gas Production from Shale Formations**

by

Daniel Zavala Araiza, B. S.

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Dedication

For Yazmin, my soulmate.

“So, these things –formerly mute and said to be objective because passive like the slaves of an activity that would belong only to us - these coding-coded things, as though awakened, speak just as much and perhaps better than us, they also say, write, sing, communicate among themselves, through a kind of reciprocal encoding, a kind of common language, a kind of music, harmonic, disharmonic.”

- *Michel Serres, Biogea*

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Atmospheric Emissions and Air Quality Impacts of Natural Gas Production from Shale Formations

Daniel Zavala Araiza, Ph.D.

The University of Texas at Austin, 2014

Supervisor: David Allen

Natural gas is at the core of the energy supply and security debates; new extraction technologies, such as horizontal drilling and hydraulic fracturing, have expanded natural gas production. As with any energy system, however, natural gas has an environmental footprint and this thesis examines the air quality impacts of natural gas production.

Greenhouse gas (GHG), criteria pollutant, and toxic emissions from natural gas production have been subject to a great amount of uncertainty, largely due to limited measurements of emission rates from key sources. This thesis reports direct and indirect measurements of emissions, assessing the spatial and temporal distributions of emissions, as well as the role of very high emitting wells and high emitting sources in determining national emissions. Direct measurements are used to identify, characterize and classify the most important sources of continuous and episodic emissions, and to analyze mitigation opportunities. Methods are proposed and demonstrated for reconciling these

direct measurements of emissions from sources with measurements of ambient concentrations.

Collectively, the direct source measurements, and analyses of ambient air pollutant measurements in natural gas production regions reported in this work improve the estimation, characterization, and methods for monitoring air quality implications of shale gas production.

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1 INTRODUCTION

1.1 Importance of Natural Gas Production from Shale Formations

New technologies have revolutionized natural gas production in the United States. This gas boom is based on the extraction of natural gas from shale formations, which have very low permeability, requiring more intensive drilling and the use of horizontal drilling and hydraulic fracturing production techniques. While conventional natural gas resources in the United States are declining, the extraction of shale gas and unconventional gas may allow the United States to become a net energy exporter [1].

It is expected that by the year 2035, 49% of U.S natural gas production will come from shale gas, with total natural gas production increasing by roughly 30% [2]. Figure 1-1 shows the geographical location of shale plays in the United States. There are important geological differences between plays, leading to differences in extraction and production methods used in each of the regions.

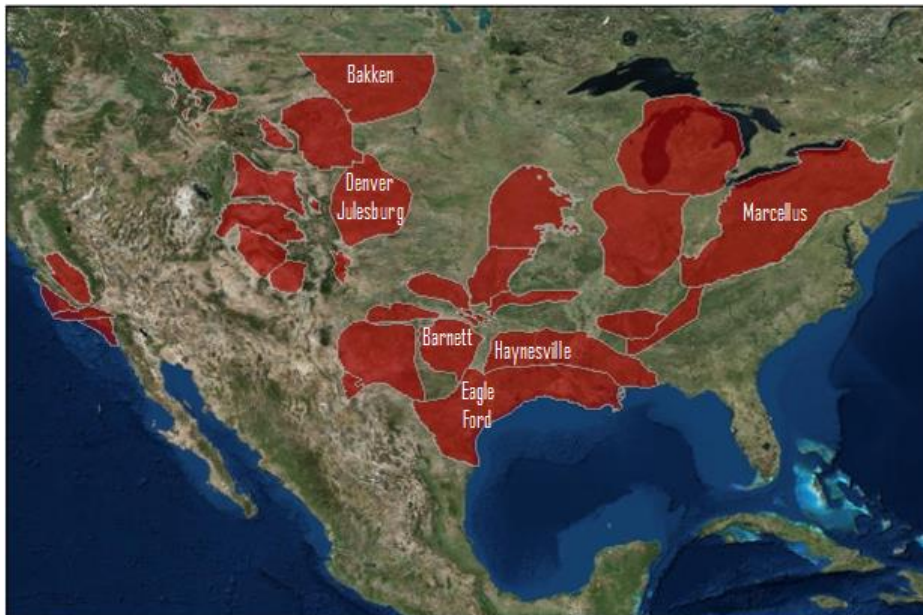


Figure 1-1. Geographical location of shale plays in the United States. (Adapted from EIA, Annual Energy Outlook [2].)

Of the various shale plays in the United States, the Barnett Shale (the focus of much of the work reported here), located in North Central Texas has among the longest histories of production. Over the course of a decade, production increased from 0.11 billion cubic feet per day (bcf/d) in 2000, to 5 bcf/d by 2011 [3-4], making the Barnett Shale one of the largest shale gas production regions in the U.S.. In comparison, the Haynesville region currently produces 7 bcf/d, the Marcellus produces 4.5 bcf/d, and Fayetteville produces 3 bcf/d [5]. Total U.S. natural gas production is approximately 75 bcf/d.

The technological developments that allow the extraction of natural gas from shale formations have been mainly developed and used in the United States, nevertheless, shale resources around the world are vast and a number of countries have started to explore the possibility of extracting shale gas and shale oil. An example of this is Mexico, where recent policy reforms have opened the door to assessment and development of national resources [6-7]. Figure 1-2 shows the ten countries with the largest estimated technically recoverable shale gas resources. The country with the most extensive Shale Gas resources is China (1,115 Trillion cubic feet (Tcf)), followed by Argentina (802 Tcf), and Algeria (707 Tcf)[8].

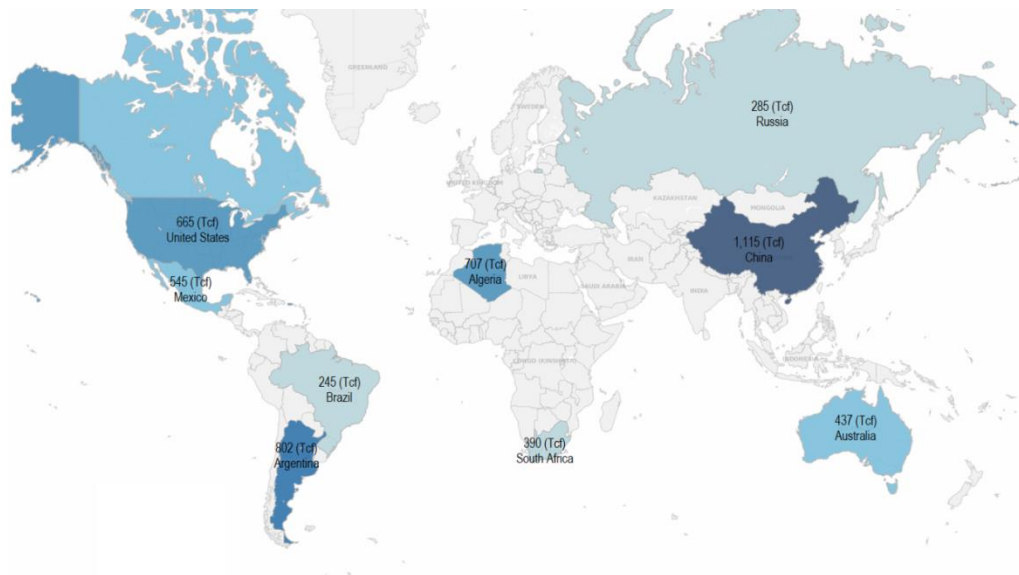


Figure 1-2. Technically recoverable shale gas for the ten countries with the biggest shale gas resources (reported in Trillion cubic feet) (Adapted from [8]).

Melikoglu [9] reported ratios of $[total\ estimated\ natural\ gas\ consumption\ (between\ 2010\ and\ 2040)]/[natural\ gas\ reserves]$ and $[total\ estimated\ natural\ gas\ consumption\ (between\ 2010\ and\ 2040)]/[natural\ gas\ reserves\ +\ shale\ gas\ resources]$ for the ten countries with the largest technically recoverable shale gas resources. For China, when only conventional natural gas reserves are considered (no shale gas resources), the ratio is 2.61, when shale gas resources are considered, the ratio decreases to 0.23. Similarly, for the United States the ratio without considering shale gas resources is 2.50, once shale gas resources are added, it decreases to 0.83. These results highlight the importance of shale gas production, and illustrate why shale gas will play a key role in meeting worldwide demand for natural gas.

The widespread availability of natural gas, both worldwide and in the United States, has caused a displacement of other fossil fuels, reshaping the overall energy landscape. In the United States, 34% of natural gas consumption is used for electricity generation, with forecasts showing that by the year 2035, natural gas will displace coal as the major source of electricity generation [10]. While changes in natural gas production are already reshaping electricity generation, these changes will also influence other sectors.

Low natural gas prices coupled with an increased supply of natural gas co-products, such as ethane, propane, butane, and natural gas liquids, will re-shape the chemical manufacturing sector, with natural gas liquids potentially replacing petroleum-derived feedstocks [10]. Production of ammonia, nitrogen based fertilizers, and hydrogen (produced by steam reforming of methane), are other examples of industrial processes that rely on methane from natural gas that will potentially be affected due to the enhanced availability of natural gas co-products and the decrease in price of natural gas.

The magnitude and rate at which shale gas production is transforming the United States, both in economic and industrial terms, makes it important to study, understand, and assess the potential environmental impacts attached to such transformations. In addition, the present state of shale production development in the United States; characterized by

decades of technological development and an advanced stage of production and extraction in many of the shale basins, can provide lessons for other countries where shale production is in an early stage, or will become a reality in the near future.

1.2 Environmental Impacts of Natural Gas Production from Shale Formations

There have been a variety of analyses of the environmental implications of shale gas production, which have addressed (i) *ground water contamination, waste water generation and water availability* [11-14], (ii) *greenhouse gas (GHG)* [15 – 18] *criteria pollutants, and air toxics emissions* [19-20], (iii) *increased seismic activity* [21-22], and (iv) *land use/ land change issues*[22, 23].

This work will focus on the air quality implications of the production of natural gas from shale formations, particularly focusing on greenhouse gases, photochemical air pollutants and their precursors, and air toxics.

Estimating the emissions of air pollutants from natural gas production is a challenging task, due to the high density of sources, the regional characteristics of the geological formations, and the wide range of spatial and temporal scales over which emissions occur. For GHGs, the emissions have impacts on national and global scales; for air toxics and criteria pollutants, the spatial scale is local and regional. Temporal scales of air quality impacts range from hours to decades. This thesis describes a set of multi-level, multi-scale analyses that can be applied to characterize air quality impacts of shale gas production.

1.3 Research Objectives

The goals of this work are to:

1. Measure and estimate GHG emissions related to shale gas production activities.
2. Measure and estimate emissions of criteria and toxic pollutants related to shale gas production.
3. Identify and analyze the distribution of emissions from shale gas production, focusing on temporal and spatial patterns.
4. Assess causal factors and emission reduction opportunities.
5. Assess the performance of current emission inventories in predicting observed emissions.
6. Develop tools for the reconciliation of measurements, at multiple spatial scales, and emission inventories

Some of the analyses will be national in scope. Others will focus on the Barnett Shale region in North Central Texas.

1.4 Dissertation Structure

Chapter 2 of this work reviews existing literature on air quality implications of natural gas production from shale gas formations. The review covers greenhouse gases, volatile organic compounds (VOC) (photochemical air pollutants), and air toxics; and identifies uncertainties in current estimates. **Chapter 3** and **Chapter 4** describe measurements done on two of the main sources of emissions from natural gas production: *pneumatic controllers* and *liquids unloading*, assessing the spatial and temporal distributions of emissions as well as the role of very high emitting wells and controllers in determining national emissions.

An important consideration when the environmental footprint of natural gas is estimated and compared against other fuels, is to acknowledge that along with natural gas, other co-products such as Natural Gas Liquids and oil are produced by gas wells. **Chapter 5** describes an allocation method for methane (greenhouse gas) emissions from natural gas production among the different co-products.; consequences and implications of the allocation scheme are discussed in the chapter.

Chapter 6 focuses on criteria pollutants and air toxic emissions. For this particular analysis, attention is shifted from the national scale, to the Barnett Shale production region in North Central Texas. The work uses dispersion modeling tools to predict ambient concentrations (based on an emissions inventory), which are compared to ambient measurements, allowing a reconciliation between top-down (ambient) measurements and bottom-up emission estimates, using VOC emissions from shale gas production as a case study.

As a follow-up to the work presented in Chapter 6; **Chapter 7** develops a methane ethane, and propane emission inventory for the Barnett Shale; based on the VOC inventory and a recently published dataset of direct measurements of methane emissions from natural gas production. The chapter discusses the spatial distribution of emissions and temporal changes in the magnitude of emissions as a consequence of changes in production over time. Based on the combination of tools developed and used for the study of GHG emissions, criteria pollutants, and air toxics, **Chapter 8** proposes a design of a multi-scale system that allows the detection and attribution of methane in natural gas production region.

Finally, **Chapter 9** discusses implications of the results, summarizing major contributions and findings. This last chapter also provides recommendations that could be explored in future work.

1.1 References

- (1) IEA. (2012). *World Energy Outlook*. Retrieved 5 29, 2013, from <http://www.worldenergyoutlook.org/publications/weo-2012/>
- (2) EIA. (2012). *Annual Energy Outlook*. DOE/EIA.
- (3) Medlock, K. B. (2012). Modeling the implications of expanded US shale gas production. *Energy Strategy Reviews*, 1, 33-41.
- (4) Rail Road Commission of Texas . (2012). *Newark, East (Barnett Shale) Statistics*. Retrieved 12 18, 2012, from <http://www.rrc.state.tx.us/data/fielddata/barnettshale.pdf>
- (5) EIA. (2013). *Annual Energy Outlook 2013*. Retrieved 10 19, 2013, from http://www.eia.gov/energy_in_brief/data/shaleplay_production_AEO2013ER.xlsx
- (6) Lozano Maya, J. (2013). The United States experience as a reference of success for shale gas development: The case of Mexico. *Energy Policy*, 70-78.
- (7) SENER (Department of Energy, Mexico). (n.d.). *Whay is Shale Gas/Oil and why is it important?* Retrieved 6 2014, from http://www.energia.gob.mx/webSener/shale/shale_en.html
- (8) EIA. (2013). *Technically Recoverable Shale Oil and Shale Gas Resources: An Assessment of 137 Shale Formations in 41 Countries Outside the United States*. Retrieved June 13, 2013, from <http://www.eia.gov/analysis/studies/worldshalegas/>
- (9) Melikoglu, M. (2014). Shale Gas: Analysis of its role in the global market. *Renewable and Sustainable Energy Reviews*, 460-468.
- (10) EIA. (2014). *Annual Energy Outlook*. Washington DC: Department of Energy.
- (11) EPA. (2011). *Plan to Study the Potential Impacts of Hydraulic Fracturing on Drinking Water Resources*. Retrieved 8 15, 2012, from http://www.epa.gov/hfstudy/HF_Study__Plan_110211_FINAL_508.pdf

- (12) Osborn, S. G., Vengosh, A., Warner, N. R., & Jackson, R. B. (2011). Methane contamination of drinking water accompanying gas-well drilling and hydraulic fracturing. *Proceedings of the National Academy of Science*, 108, 8172-8176.
- (13) Nicot, J., & Scanlon, B. R. (2012). Water Use for Shale-Gas Production in Texas. *Environmental Science & Technology*, 46, 3580-3586.
- (14) Rahm, B., & Riha, S. (2012). Toward strategic management of shale gas development: Regional, collective impacts on water resources. *Environmental Science & Policy*, 17, 12-23.
- (15) Alvarez, R. A., Pacala, S. W., Winebrake, J. J., Chameides, W. L., & Hamburg, S. P. (2012). Greater focus needed on methane leakage from natural gas infrastructure. *Proceedings of the National Academy of Sciences*, 109(17), 6534-6440.
- (16) Howarth, R. W., Santoro, R., & Ingraffea, A. (2011). Methane and the greenhouse-gas footprint of natural gas from shale formations. *Climatic Change*, 106, 679-690.
- (17) O'Sullivan, F., & Paltsev, S. (2012). Shale gas production: potential versus actual greenhouse gas emissions. *Environmental Research Letters*, 7, 1-7.
- (18) Wigley, T. L. (2011). Coal to gas: the influence of methane leakage. *Climatic Change*, 108, 601-608.
- (19) Armendariz, A. (2009). *Emissions from natural gas production in the Barnett shale area and opportunities for cost-effective improvements*. Austin, Texas: Environmental Defense Fund.
- (20) Rahm, D. (2011). Regulating hydraulic fracturing in the shale gas plays: The case of Texas. *Energy Policy*, 39, 2974-2981.
- (21) Ellsworth, W. L. (2013). Injection-Induced Earthquakes. *Science*, 341.

(22) Davies, R., Foulger, G., Bindley, A., & Styles, P. (2013). Induced seismicity and hydraulic fracturing for the recovery of hydrocarbons. *Marine and Petroleum Geology*, 45, 171-185.

(23) Jenner, S., & Lamadrid, A. J. (2013). Shale gas vs. coal: Policy implications from environmental impact comparisons of shale gas, conventional gas, and coal on air, water, and land in the United States. *Energy Policy*, 53, 442-453.

2 BACKGROUND

The previous Chapter discussed the importance of natural gas, and how the development of new technologies has enabled an expansion in the production of unconventional resources, such as shale gas. The environmental impacts associated with shale gas production have made it controversial. This Chapter reviews the impacts associated with greenhouse gas emissions, criteria pollutants, and air toxics. The potential carbon footprint of natural gas production is described, with a focus on the assumptions embedded in its calculation. Existing datasets of GHG emissions, criteria pollutants, and air toxics are presented.

The present state of knowledge of the air quality implications of shale gas production is characterized by great uncertainties in emission estimates. This chapter also discusses the uncertainties and describes how recent research efforts have begun the process of improving understanding of these emissions.

2.1 Greenhouse Gas Emissions from Shale Gas Production

2.1.1 *Carbon footprint of Natural Gas*

Greenhouse gas (GHG) emissions from the combustion of natural gas are lower (per unit of energy) than for other fossil fuels used for the production of energy, such as coal and petroleum.

To generate a MJ of energy, 19.9 g of methane, the primary component of natural gas, must be combusted. If complete combustion is assumed, 54.7 g of carbon dioxide would be released. In contrast, 44 g of coal would be needed to generate the same MJ of energy¹, and if the coal has a composition of 85% carbon, its combustion would yield 137 g of carbon dioxide. Thus, if the analysis is based on the emissions released as a

¹ Assuming a heating value of 50.3 MJ/kg for methane and of 22.7 MJ/kg for coal (lower heating values) [1-2].

consequence of the combustion of fossil fuels for the production of energy, natural gas has considerably lower carbon dioxide emissions than coal (and petroleum) (Figure 2-1).

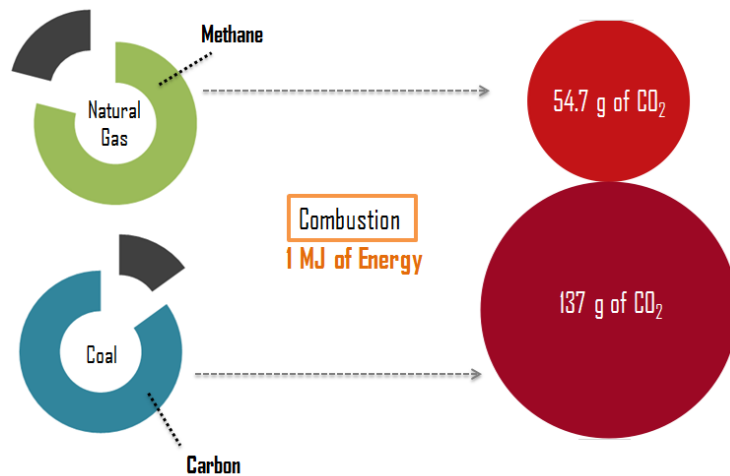


Figure 2-1. Comparison of GHG emissions from the generation of 1 MJ of energy from the combustion of methane (natural gas) and coal.

While greenhouse gas emissions from combustion of natural gas are lower than for other fossil fuels, methane, CH₄, the principal component of natural gas, is a potent GHG; its emissions along the natural gas supply chain could occur in such an extent that the lower combustion footprint observed when compared against other fossil fuels could be offset. The amount of methane emissions that would alter the GHG footprint of natural gas, relative to other fuels, depends on the potency assumed for methane.

Emissions of greenhouse gases are often expressed on the basis of the radiative forcing that they cause (in units of carbon dioxide equivalents). A number of assumptions are associated with the calculation of carbon dioxide equivalents. One of the primary assumptions is the time over which radiative forcing is integrated. Most assessments (such as those done by the Intergovernmental Panel on Climate Change) have assumed a 100 year time horizon, over which radiative forcing is integrated and averaged. For methane, which is converted to CO₂ in the atmosphere, over decadal time scales, the choice of averaging time is important in establishing the importance of the emissions. If a 100 year time horizon is used, methane has a radiative forcing 34 times higher than CO₂ on a mass basis (1 kg CH₄ is equivalent to 34 kg CO₂). If a 20 year time horizon is used,

the radiative forcing is 86 times higher than CO₂. The radiative forcing is 102 times the value for CO₂ if immediate effects are considered [3]. Figure 2-2 shows how the contributions to radiative forcing from the 2012 U.S. national inventory of greenhouse gas emissions [4] would appear using different time horizons, for the three GHG with most emissions (CO₂, CH₄, and N₂O)².

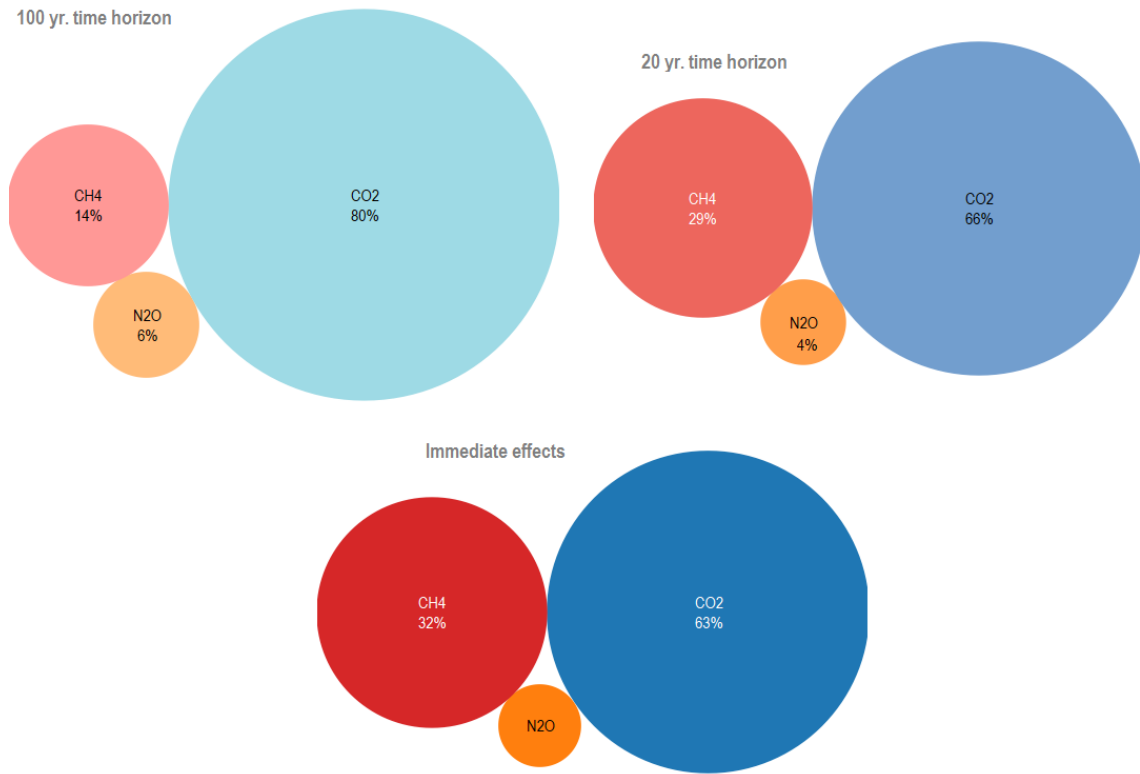


Figure 2-2. Total GHG Emissions by species from the EPA GHG National Inventory for 2012 [4]. The circles are sized relative to the percentage of total emissions considering the Global Warming Potentials (GWP) for 100 year, 20 year, and immediate effects time-horizon.

² The EPA used a methane radiative forcing of 21 (100-yr time horizon) for the 2012 GHG inventory [4] (consistent with the IPCC fourth assessment report (AR4) [5]). The IPCC AR5 (released in 2013) updates this value to a radiative forcing of 34 [3].

Thus, methane may represent 15% to a third of radiative forcing due to greenhouse gas emissions in the United States, depending on whether the concern is immediate impacts, impacts over a 20 year time horizon or impacts over a century. The climatic implications of transitioning to a higher consumption and usage of natural gas will depend on the selected time horizon, but by any measure, the potential impacts could be large.

Alvarez, *et al* [6], performed a series of analyses comparing greenhouse gas emissions associated with the use of natural gas to other fossil fuels, assuming a variety of GWP. Their study highlighted that, depending on the assumed GWP (and selected time horizon), for some fuel-switching scenarios, net climate benefits would require methane leak rates of less than 1%, while for other scenarios, with long term climate benefits, the leak rate would be set at 4% or more. This work frames the range of potential methane leak rates that would produce positive climate benefits as a result of replacing other fuels with natural gas. However, the study makes clear the need for accurate estimation and measurement of emissions.

2.1.2 Methane Emissions along the Natural Gas Supply Chain

Based on the EPA National GHG inventory for 2012 6,186 Gg of methane were released from natural gas systems [4], This corresponds to 210.3 Tg of CO₂ equivalents (Eq.), using a 100-year time horizon (Methane Global Warming Potential (GWP) of 34); However, if the 20-year time horizon is selected (Methane GWP of 86), methane emissions from natural gas systems would result in 532.0 Tg of CO₂ Eq. and if the immediate effects scenario (Methane GWP of 102) is used, methane emissions from natural gas systems would be 631.0 Tg of CO₂ Eq.

The emissions of methane can be attributed to various parts of the natural gas supply chain, and the supply chain can be divided into four main stages: production, processing, transmission, and distribution (Figure 2-3). During the *production stage*, natural gas, with a mixture of hydrocarbon liquids and water, are extracted from wells. The mixture is typically sent to a separator unit, removing gas from the top of the separator and sending

water and hydrocarbon liquids (condensate) to storage tanks. The production stage, including well operation as well as pre-production activities (e.g., well completion), is estimated to account for 32% of GHG emissions from natural gas systems in the 2012 EPA inventory [4]. The *Processing* stage involves the removal of various constituents in order to produce “pipeline quality” gas. This stage is estimated to account for 14% of natural gas system emissions in the 2012 EPA inventory. *Transmission* involves the flow of gas from the production sites into pipelines and storage reservoirs. The transmission stage is estimated to account for 33% of the GHG emissions in the 2012 inventory for natural gas systems. Finally, the *distribution* stage involves the delivery of natural gas to the final consumer, and is estimated to account for 20% of the 2012 GHG emissions for natural gas systems.

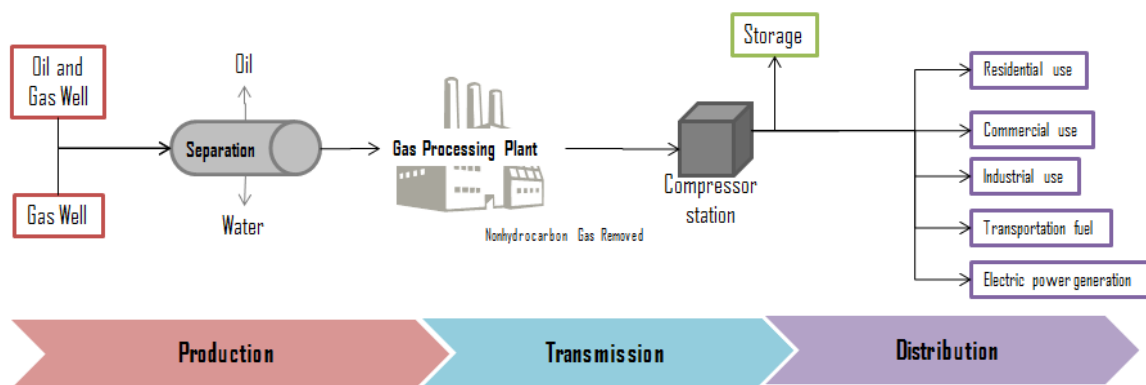
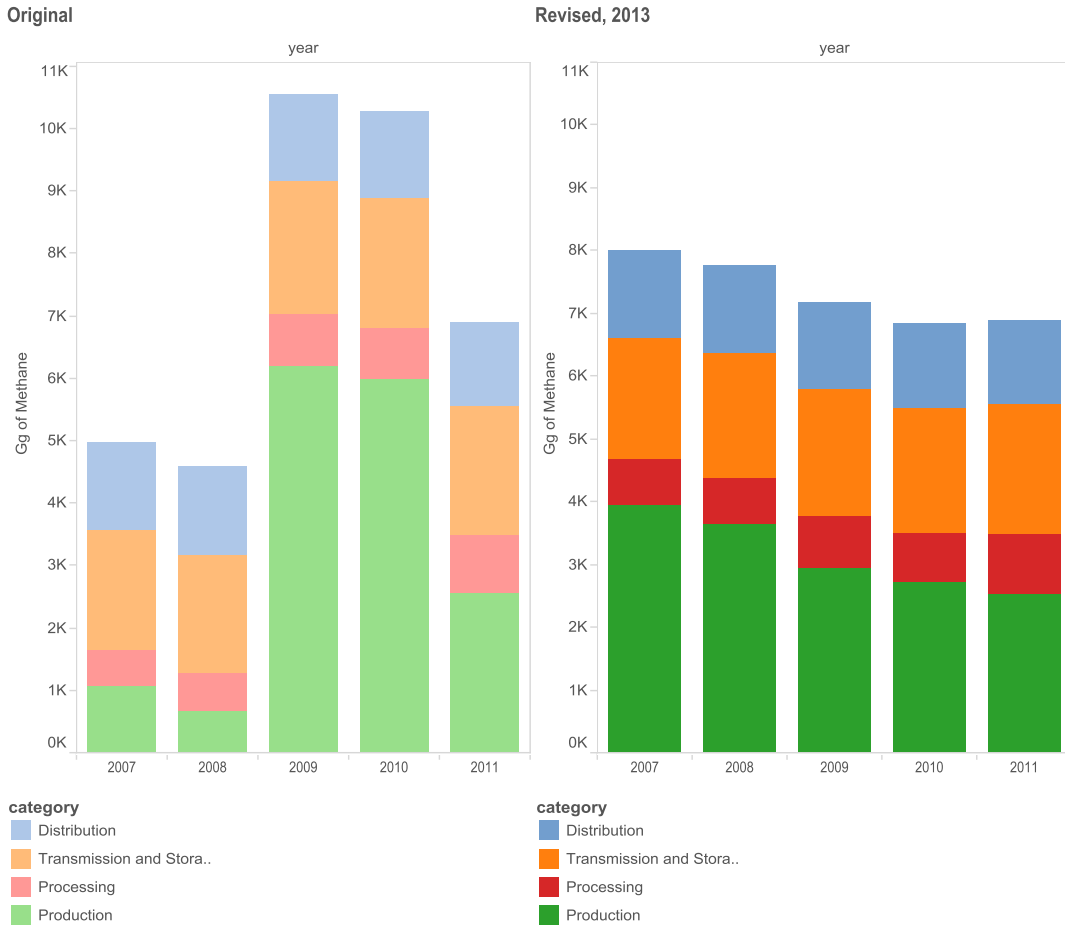


Figure 2-3. Simplified diagram illustrating the different stages in the Supply Chain of Natural Gas (Adapted from [7]).

There is considerable uncertainty in the methane emissions reported in EPA national inventories. During the past five years, GHG emissions estimated in the national inventory for natural gas production have varied by an order of magnitude. Figure 2-4 reflects these changes [8]. This variation has been caused mainly by significant modifications in the estimation methods. When the EPA national inventories for the years 2008 and 2009 were originally published, GHG emissions from the production stage were estimated as 674 and 6,205 Gg of methane, respectively. If the emissions are

estimated using a consistent methodology, changes from year to year are much smaller (Figure 2-4, right).



(a) **(b)**
Figure 2-4. CH₄ Emissions from Natural Gas Systems category in the EPA National Emission inventory; comparing (a) the year when the inventory was originally published against (b) the values reported in 2013 after revisions made to the estimation methods were applied consistently to previous years [8].

Overall, methane leak rates in the natural gas supply chain (methane as a volume percentage of natural gas production with all methane emissions allocated to natural gas) reported in EPA inventories over the past several years have ranged from less than 1% in

2008 to 2.1% in 2009, with a current estimate of 1.3%. Other researchers have suggested that the leak rates may be much higher than 2%, with some values as high as 8% or more [9]. The Department of Energy and Climate Change in the United Kingdom prepared a review of available estimates of greenhouse gas emissions from shale gas production [10], based exclusively on United States data. Similarly, Caulton, *et al* [11] report ranges of published methane emission estimates; with a range of 0.6 – 7.7% of the lifetime production for the *upstream* and *midstream* sector, and 0.07 to 10% emitted during the downstream stage. These publications document the wide range of published estimates and emphasize the great uncertainty among greenhouse emission estimates.

The production stage represents the sector in which the greatest changes have occurred in the EPA inventory. Table 2-1 summarizes emissions reported in the EPA national emission inventory from 2011. In assembling the national inventory, the EPA estimates potential emissions from source categories, and then reduces the potential emissions by estimated voluntary reductions, as well as reductions required by regulations.

Table 2-1. EPA National emission inventory estimates by source category (Adapted from [8]).

Environmental Protection Agency source activity	Potential emissions (Gg)	Emission reductions (Gg)	Net Emissions (Gg)
Completions with hydraulic fracturing	1,221	567	654
Refractures (workovers with hydraulic fracturing)	266	124	143
Pneumatic device vents	1,134	779	355
Chemical injection pumps	64	30	34
Equipment leaks: gas wells	52	24	
Equipment leaks: separators	107	50	
Equipment leaks: meters/piping	102	48	172
Equipment leaks: heaters	33	15	
Equipment leaks: dehydrators	31	15	
Workovers without hydraulic fracturing	0.6	0.3	0.3
Liquids unloading	257	0	257
Kimray pumps	365	180	
Condensate tanks	313	167	
Gas engines	276	49	
Dehydrators vents	114	73	
Reciprocating compressors	84	35	
Pipeline leaks	170	80	
Well drilling	0.8	0.4	
Blowdowns	6.7	2.3	930
Compressor starts	6	3	
Pressure relief valves	0.7	0.3	
Mishaps	2	1	
Emissions from coal bed methane and offshore production			
Coal bed methane–produced water	59	27	
Offshore and deepwater platforms	289	136	
TOTAL	4,949	2,405	2,545

The source category with the highest reported emissions in 2011 was completion flowbacks (the 2012 emission inventory, released in 2014, reports a substantial reduction in these emission, largely based on the measurements reported in this work); three other

important source categories are pneumatic devices, equipment leaks and liquid unloadings (detailed description of these categories is provided in *section 2.1.3*).

Part of the reason for emission estimation uncertainty is a lack of measurement data on the methane emissions from natural gas production activities. To address this lack of data, a team led by the University of Texas collected data on a variety of natural gas production activities during 2012. Data were collected on completion flowbacks, wells in routine production, and liquid unloadings. Descriptions of each of these sources and results from the measurements are described in the next section.

2.1.3 Direct Measurements of Methane Emissions at Natural Gas Production Sites

To address the lack of data on methane emissions from natural gas production, a team led by the University of Texas collected data on a variety of natural gas production activities during 2012 [12]. Data were collected on completion flowbacks, wells in routine production, and liquid unloadings.

Allen, *et al.* [12] reported measurements at roughly 478 gas wells; if it is assumed that the measurements are representative of the national population, methane emissions from the source categories on which measurements were made would sum to 957 Gg of methane, which can be compared against the 1,200 Gg of methane reported in the EPA national inventory for comparable source categories [8]. While in total, the emission measurements agreed with the total emissions estimated in the inventory, some source categories had much higher emissions than expected; other source categories had much lower emissions than expected. .

Allen, *et al.* found emissions from well completions to be 97% lower than the 2011 EPA emission estimates (released in 2013) [8]. Pneumatic devices and equipment leaks were found to account for 40% of the methane net emissions from natural gas production, with measured emissions being 70% and 50% higher than the EPA net emission estimates, respectively. Allen, *et al.* acknowledged the need for additional measurements to

characterize and understand the distribution of emissions from pneumatic devices and liquids unloading.

Well completion flowbacks: Once a well is drilled and the fracturing has taken place, the well must be cleaned of sand and fluids that were previously injected; the recovery of these fluids is known as flowback. During this stage, natural gas dissolved or entrained in the flowback liquids can be captured, controlled or emitted. The EPA national GHG emissions inventory reported a total of 8,077 completion flowback events during 2011, which accounted for 26% of the emissions related to natural gas production [8].

Well completion flowback emissions reported by Allen, *et al.* were the first measurements from this source to be reported in the scientific literature. From the 27 wells completions that were sampled, flowback durations ranged from 5 hours to 339 hours, with an average emission of 1.7 Mg of methane, a minimum of 0.01 Mg and a maximum of 17 Mg (which corresponds to the completion that lasted 339 hours).

Potential emissions are defined as methane leaving the wellhead; which may or may not be sent to an emissions control device. From the 27 completion events sampled by Allen *et al*, the average potential emissions were 124 Mg, however, 67% of the flowbacks had emission control devices, and 33% had no capture or control Allen, *et al* [12]. The average control efficiency $((1 - (\text{methane emitted to the atmosphere} / \text{potential emissions})) * 100)$ was 99%. Figure 2-5 summarizes data from the 27 completion events that were directly measured. The 9 events where no emission control devices were in place are represented by the darker green bars; where the measured emissions equal potential emissions. The highest emitting events had actual measured emissions representing 1% of potential emissions for those wells.

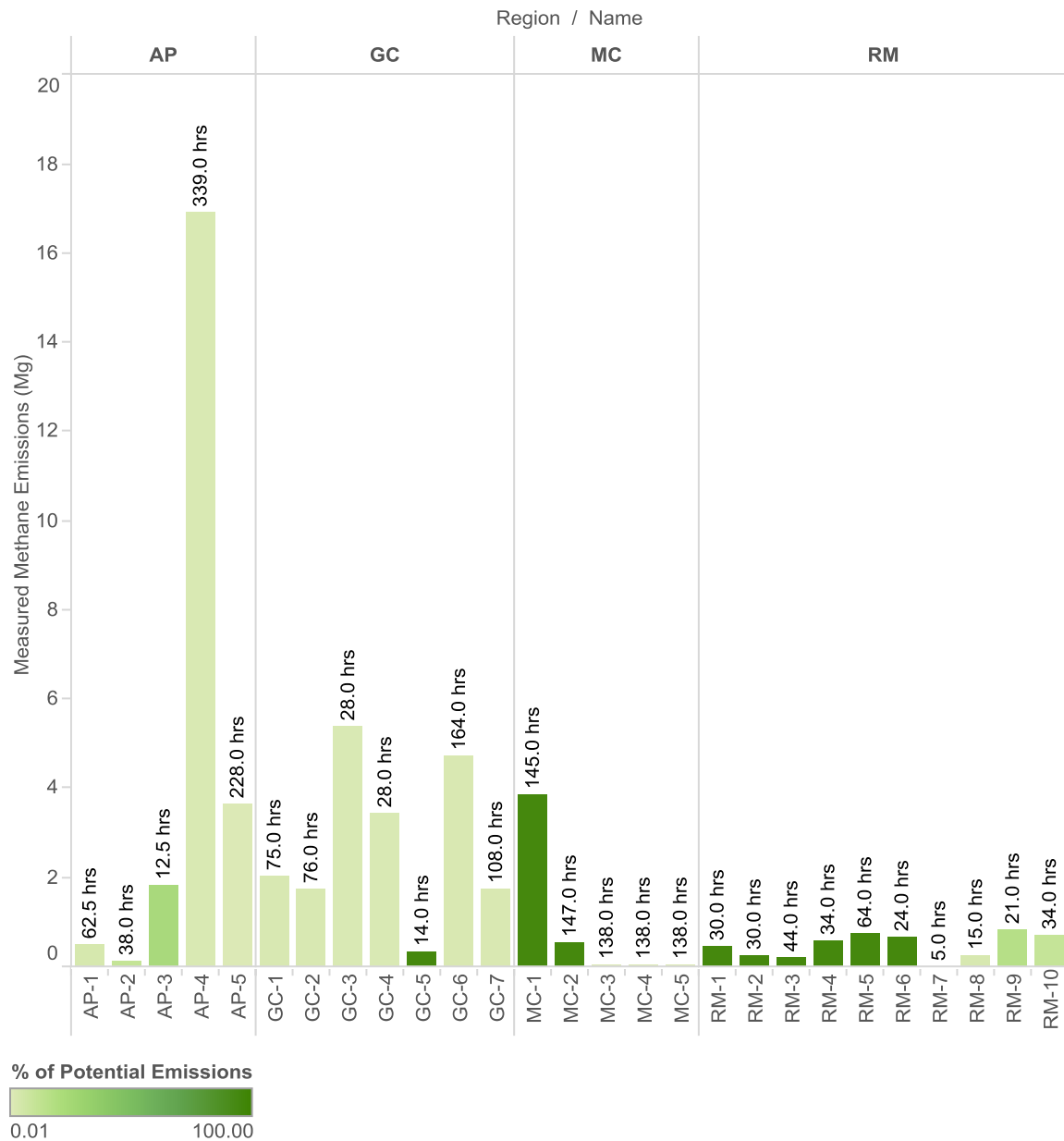


Figure 2-5. Well completion events sampled by Allen et al. The height of the bars represents the total methane emissions measured in each event. The color represents the percentage of potential emissions that actual measured emissions represent. The duration of the event is shown at the top of each bar. Regions are designated as AP= Appalachians, GC= Gulf Coast, MC= Midcontinent, RM= Rocky Mountains.

Wells in routine production: Once wells go into routine production, the operation of pneumatic controllers, chemical injection pumps, tanks, and compressors, along with equipment leaks, lead to methane emissions. These sources account for 32% of the emissions reported in the EPA national inventory for the natural gas production sector [8]. Among the sources related to routine production, pneumatic controllers stand out due to high equipment counts and heterogeneity of design and operation.

Pneumatic controllers: In the work reported by Allen, *et al.*[12], a total of 305 pneumatic controllers were sampled with an average emission rate of 10.5 scf/h of methane (11.2 scf.h of whole gas), a median value of 3.68 scf/h, and a maximum value of 126.9 scf/h. The emissions from pneumatic controllers were the largest single contributor to methane emissions from natural gas production. The data showed significant geographical variability and variability by equipment type. The importance of this source category, and the high degree of variability, led to the decision to collect additional data. Figure 2-6 shows the distribution of pneumatic controllers by region and by host company, and also highlights the impact of outliers on the emission average. Chapter 3 discusses the findings of a second stage of measurements that specifically targeted pneumatic controllers.

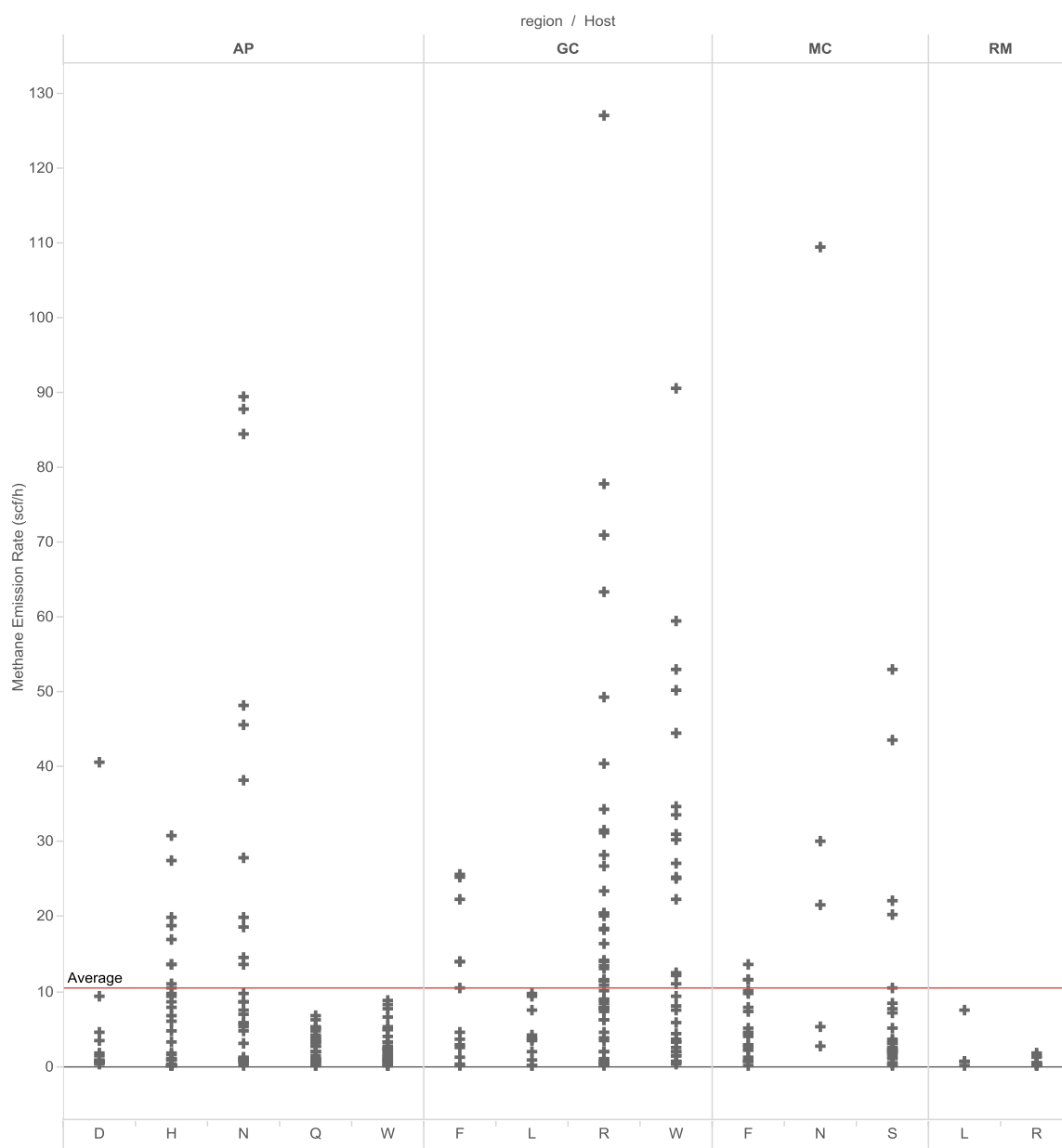


Figure 2-6. Distribution of methane emission rates from pneumatic controllers sampled by Allen, et al. The devices are categorized by region (AP= Appalachians, GC= Gulf Coast, MC= Mid Continent, and RM= Rocky Mountains) and host company.

Liquids unloadings: When the velocity of natural gas up the subsurface tubing in a well declines and is not enough to lift liquids that are co-produced with the gas, liquids accumulate in the wellbore, restricting the flow of gas, and the accumulated liquids must

be removed. When a *liquids unloading* takes place, flow is diverted into an atmospheric pressure tank, causing a pressure gradient that increases the velocity of the fluid. If there is no emissions control system in place, gas is emitted to the atmosphere. In the 2011 national inventory, EPA estimated that unloading events accounting for 10% of methane emissions from natural gas production [8].

Direct methane measurements conducted by Allen *et al.* [12] took place in four different production regions throughout the United States: Appalachia, Gulf Coast, Midcontinent, and Rocky Mountains. Direct measurements were made at 150 production sites (489 wells), 27 well completion flowbacks, and 9 well unloadings. The nine unloadings measurements represented the first measurements of liquid unloading emissions reported in the scientific literature. These measurements are the starting point for the work presented in Chapter 4, which analyses in detail a second stage of direct measurements that target liquids unloading events.

The nature of the unloading events sampled by Allen *et al.* [12] was diverse, with episodes lasting from 10-15 minutes of uninterrupted flow to two hours of uninterrupted flow. The average measured methane emission was 1.1 Mg per event; with a minimum value of 0.02 Mg and a maximum value of 3.67 Mg of methane. Figure 2-7 shows the measured methane emissions of each measured event and the frequency of liquid unloadings per year for that particular well.

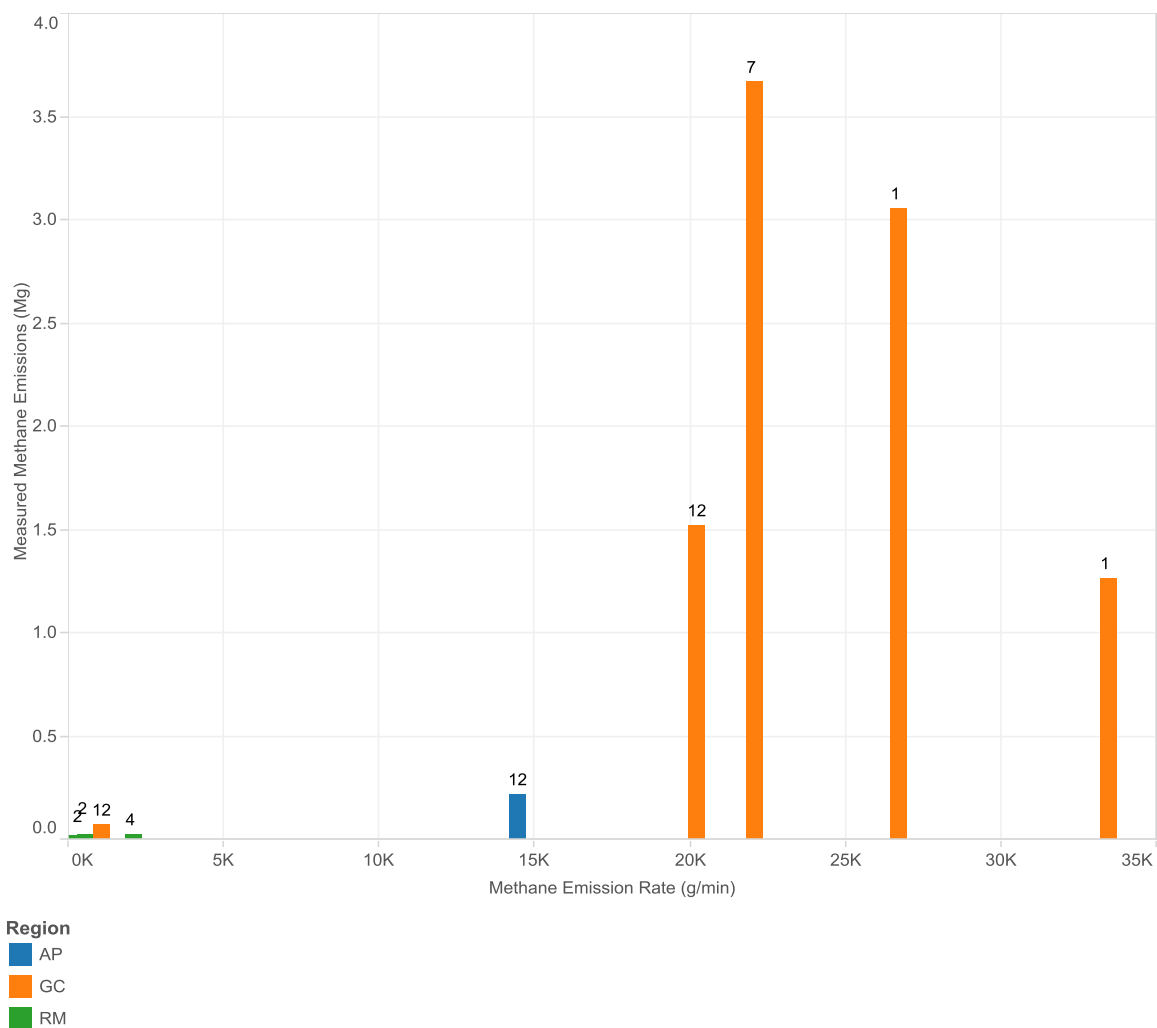


Figure 2-7. Methane emissions from the 9 liquid unloadings directly measured by Allen et al. The plot shows the methane emission rate for each event, as well as the number of events per year per well at the top of each bar, and the region where the event took place (AP= Appalachians, GC= Gulf Coast, RM= Rocky Mountains).

The diversity of these events and the small sample size makes it difficult to extrapolate and compare to national inventories

2.1.4 Comparison among measurements and emission estimates

Ambient measurements made at ground level and aloft using aircraft can also be used to improve understanding of emissions from natural gas production.

Ground level measurements, made downwind of natural gas sites, have made use of a variety of techniques designed to infer emission rates from ambient concentrations. One of the most precise methods involves the release of tracer compounds (e.g., SF₆, N₂O, and C₂H₂) at a known rate at or near the methane emission source [12, 13-17]. Downwind measurements of methane (minus background) and the tracers (minus background) are equal to the ratio of emission rates if the dispersions of the methane and the tracer are identical. Consequently, methane emissions can be estimated by multiplying the known emission rate of the tracer by the background corrected downwind concentration ratio of methane to the tracer. Several of these studies have pointed to a skewed distribution of emissions among sites, with a small number of sites accounting for a large fraction of emissions. The existence of high emitting sources (e.g., a pneumatic controller valve that emits much more than the average pneumatic controller, or a liquid unloading event) may explain some of this variability, [12,18].

However, a careful interpretation of these skewed distributions is needed. For example, in a study conducted for the City of Fort Worth [19], emissions from 375 well sites in the Barnett Shale production region were reported (sites were randomly selected from the well sites that were within the City of Fort Worth); 30% of the sites had one well, 63% had between 2 and 6 wells, and one site had 13 wells. Similarly, whereas 78% of the sites had between 1 and 4 tanks, 16% had more than 4 tanks, and one site had 20 tanks. The potential sources of fugitive emissions, such as valves and flanges, varied by an order of magnitude or more between sites. Ten percent of the sites had less than 62 valves, but 10% had more than 446 valves. Ten percent of the sites had 390 or less connectors (such as flanges), but 10% had more than 3,571. Because of this heterogeneity in the equipment among sites, simple comparisons of methane emissions among sites, without adjustments for equipment counts, should be viewed with caution.

Ambient measurements made by aircraft have also been used to assess emissions from the natural gas supply chain. In these studies, average aloft concentrations, upwind and downwind of a natural gas production region, are determined. To estimate total emissions for a region, the difference between upwind and downwind concentration is multiplied by the advection rate of air over the basin (mixing height multiplied by the average wind velocity and the horizontal dimension of the basin). In order to estimate emissions related to natural gas operations, emission from the rest of the sources (e.g emissions from livestock and landfills) must be estimated and subtracted from the total emissions from the area. The accuracy and assumptions made to estimate emissions of non-natural gas sources impacts the uncertainty of aircraft based measurements.

In some regions, aircraft based emission estimates have been a very high percentage of the region's natural gas production or use. Aircraft measurements were reported for a natural gas and oil field in Utah, with an estimated emission rate in the range of 6.2 – 11.7% of natural gas production. In aircraft-based analyses of methane emissions in Los Angeles, fugitive losses from natural gas pipelines and the urban distribution system, along with geological seeps, were identified as the dominant emission sources (47% of total emissions) [20-22] and accounted for 2% of the natural gas delivered to customers, and 1% of the gas flowing into the basin [21]. These analyses suggest emissions sources that are not accurately accounted for in current emission inventories for both natural gas production and natural gas delivery systems. Again, however, these data must be interpreted with care. The attribution of methane emissions to the natural gas supply chain from regions that have both oil and gas production should be done carefully. Gas production operations, and their emissions, also vary over the life of a gas field. Emissions from wells in routine operation may stay roughly constant through the life time of the well; episodic emissions such as new well completions and liquid unloadings are dependent upon the age of the well or the gas field. Early in a field's life, drilling and well-completion activities will be more common than later in a field's life. Late in a field's life, wells may accumulate liquids, and methane venting may occur through

liquids unloading, which removes liquids from the well bore. Consider a detailed example of how liquids unloadings may skew instantaneous measurements. Allen *et al.* [12] have observed emission rates for single liquids unloading events that ranged from roughly 100 g per min to more than 30,000 g per min. These rates are much higher than emission rates for production sites (typically tens of g of methane per minute per well) or from completions (typically a few hundred g per event per min). At these emission rates, a single unloading event could, during the period that it is occurring, result in emissions that are the equivalent of just a few wells in routine production to the equivalent of up to several thousand wells in routine production. Because not all gas fields have wells that unload, and because gas wells may unload for only part of their production life cycle, emissions from different gas fields would be expected to vary, and an individual gas field would be expected to vary in its emissions over time. Overall, reconciliations between instantaneous ambient measurements should carefully account for the status of the wells in a gas field.

Collectively, recent measurements suggest that both ambient and direct source measurements will be important in examining emissions along the natural gas supply chain. *Chapter 6* and *Chapter 7* demonstrate methods for reconciliation between top-down and bottom-up measurements.

2.2 Criteria Pollutants and Toxic Emissions from Natural Gas Production

Measurements of photochemically generated air pollutants, such as ozone, in rural production regions, have motivated the collection of data on ozone precursors [23-25] from natural gas production activities. Both ozone and air toxics represent health risks, and as natural gas production increases, determining the magnitude and effect of ozone precursors and air toxics on air quality has become increasingly important.

Armendariz [23] estimated criteria pollutant and air toxics emissions from oil and natural gas sources in the Barnett Shale, estimating that emissions from VOC and NO_x related to

oil and gas production were 36% greater than emissions from motor vehicles in the region. Annual average VOC emissions for 2009 were estimated at 139 tons/day; with a 2009 natural gas production of 4.9 Bcf/day [26]; or 28.4 tons per Bcf of natural gas production. For the case of air toxic compounds, the study found that sources such as condensate tanks and engines could emit a significant amount of benzene and formaldehyde.

Similar studies have been published for other production regions in the U.S., Roy, *et al.* [25] developed a criteria pollutant emission inventory for the development, production, and processing stages of natural gas production in the Marcellus Shale (North East U.S.). The results were extrapolated to 2020, after accounting for the effects of existing and potential regulations. Estimated VOC emissions from gas production activities were 345 tons/year for 2009 and 100 tons/day for 2020, which would account for 12% of the total VOC emissions in the region.

Gilman, *et al.* [27] measured VOC emissions related to natural gas and oil operations in northeastern Colorado, establishing that light alkanes dominate among the VOC species, and that the source signature is clearly differentiated from urban sources. Their results also showed that reactivity of VOC is driven by large abundance of relatively low reactive species (light alkanes). Also in the Rocky Mountain region, Edwards, *et al.* [28] focused on the Uintah Basin in Utah, a region with increasing oil and gas production, where unusual wintertime ozone episodes (where ozone concentrations were above levels harmful to human health) have been present. The authors examine the chemical and physical processes that would lead to elevated ozone concentrations during wintertime, as a result of increased emissions from the natural gas and oil production activities in the basin.

In this thesis, the emissions of Volatile Organic Compounds (VOC) (ozone precursors) from shale gas production are examined using emission inventories and air quality monitoring data from the Barnett Shale production region in north central Texas. What

distinguishes this work from previous studies is the use of a long time series (20 month) of measurements, in contrast to short term, episodic studies. Chapter 6 presents the analyses done to VOC and air toxics emissions related to shale production activities.

2.3 Summary

The present state of knowledge of the air quality implications of shale gas production is characterized by a great uncertainty in emission estimates. Recently published data sets of direct measurements have reduced the uncertainty of the emissions and have identified the main sources from natural gas production. Nonetheless, additional measurements and analysis efforts are needed to continue to improve understanding of emission sources.

The following chapters describe how the research presented in this thesis contributes to reducing the uncertainty of emission estimates, characterizing emissions from the different sources, and reconciling different measurement and estimation schemes.

2.4 References

- (1) Green, D. W., & Perry, R. H. (2007). *Perry's Chemical Engineers' Handbook*. Blacklick, OH: McGraw Hill.
- (2) Argonne National Laboratory. (n.d.). The Greenhouse Gases, Regulated Emissions, and Energy Use In Transportation Model, GREET 1.8d.1. Retrieved 6 16, 2014, from GREET: <http://greet.es.anl.gov/>
- (3) IPCC. (2013). Fifth Assessment Report (AR5), Climate Change 2013: The Physical Science Basis. Retrieved 10 19, 2013, from http://www.climatechange2013.org/images/uploads/WGIAR5_WGI-12Doc2b_FinalDraft_All.pdf

- (4) EPA. (2014, April 15). Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2012. Retrieved 6 20, 2014, from <http://www.epa.gov/climatechange/ghgemissions/usinventoryreport.html>
- (5) IPCC. (2007). Fourth Assessment Report (AR4), Climate Change 2007: The Physical Science Basis.
- (6) Alvarez, R. A., Pacala, S. W., Winebrake, J. J., Chameides, W. L., & Hamburg, S. P. (2012). Greater focus needed on methane leakage from natural gas infrastructure. *Proceedings of the National Academy of Sciences*, 109(17), 6534-6440.
- (7) EIA. (2013). The Natural Gas Production, Transmission and Distribution System. Retrieved 5 29, 2013, from <http://www.window.state.tx.us/specialrpt/energy/nonrenewable/gas.php>
- (8) EPA. (2013, April 12). Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2011. Retrieved 4 22, 2013, from <http://www.epa.gov/climatechange/ghgemissions/usinventoryreport.html>
- (9) Howarth, R. W., Santoro, R., & Ingraffea, A. (2011). Methane and the greenhouse-gas footprint of natural gas from shale formations. *Climatic Change*, 106, 679-690.
- (10) MacKay, D. J., & Stone, T. J. (2013). *Potential Greenhouse Gas Emissions Associated with Shale Gas Extraction and Use*. London: DECC.
- (11) Caulton D. R., Shespon P. B., Santoro, R. L., Sparks J. P, Howarth, R. W., Ingraffea, A. R., Cambaliza M. O. L., Sweeney, C., Karion, A., Davis, K. J., Stirm, B. H., Montzla S. A., Miller, B. R. (2014). Toward a better understanding and quantification of methane emissions from shale gas development. *Proceedings of the National Academy of Sciences*, 111, 6237 – 6242.

- (12) Allen, T. D., Torres, M. V., Sullivan, D., Harrison, M., Hendler, A., Herndon, S. C., et al. (2013). Measurements of Methane Emissions at Natural Gas Production Sites. *Proceedings of the National Academy of Sciences*. 110, 17768-17773.
- (13) Lamb BK, Shorter JH, McManus JB, Kolb CE, Mosher BW, et al. (1995). Development of atmosphere tracer methods to measure methane emissions from natural gas facilities and urban areas. *Environ. Sci. Technol.* 29:1468–78
- (14) Shorter JH, McManus JB, Kolb CE, Allwine EJ, Siverson R, et al. (1997). Collection of leakage statistics in the natural gas system by tracer methods. *Environ. Sci. Technol.* 31:2012–19
- (15) Kolb CE, Herndon SC, McManus JB, Shorter JH, Zahniser MS, et al. (2004). Mobile laboratory with rapid response instruments for real-time measurements of urban and regional trace gas and particulate distributions and emission source characteristics. *Environ. Sci. Technol.* 38:5694–703
- (16) Herndon SC, Jayne JT, Zahniser MS, Worsnop DR, Knighton B, et al. (2005). Characterization of urban pollutant emission fluxes and ambient concentration distributions using a mobile laboratory with rapid response instrumentation. *Faraday Discuss.* 130:327–39
- (17) Herndon SC, Floerchinger C, Roscioli JR, Yacovitch TI, Franklin JF, et al. (2013). Measuring methane emissions from industrial and waste processing sites using the dual tracer flux ratio method. Presented at *Annu. Meet. Am. Geophys. Union*, Dec. 2013, San Francisco
- (18) Allen DT, Torres VM, Thomas J, Sullivan D, Harrison M, et al. (2013). Measurements of Methane Emissions at Natural Gas Production Sites: Study Appendices and Database. Austin: Univ. Tex. <http://dept.ceer.utexas.edu/methane/study/>

- (19) East. Res. Group/Sage Environ. Consult. (2011). City of Fort Worth natural gas air quality study: final report, July 13, Fort Worth, TX. Accessed Jan. 2013. http://fortworthtexas.gov/uploadedFiles/Gas_Wells/AirQualityStudy_final.pdf
- (20) Petron G, Frost G, Miller BR, Hirsch AI, Montzka SA, *et al.* (2012). Hydrocarbon emissions characterization in the Colorado front range: a pilot study. *J. Geophys. Res.* 117(D4):D04304
- (21) Peischl J, Ryerson TB, Brioude J, Aikin KC, Andrews AE, *et al.* (2013). Quantifying sources of methane using light alkanes in the Los Angeles Basin, California. *J. Geophys. Res. Atmos.* 118:4974–90
- (22) Karion A, Sweeney C, Petron G, Frost G, Hardesty RM, *et al.* (2013). Methane emissions estimate from airborne measurements over a western United States natural gas field. *Geophys. Res. Lett.* 40:4393–97
- (23) Armendariz, A. (2009). *Emissions from natural gas production in the Barnett shale area and opportunities for cost-effective improvements*. Austin, Texas: Environmental Defense Fund.
- (24) Adgate, J. L., Goldstein, B. D., McKenzie, L. M., (2014) Potential Public Health Hazards, Exposures and Health Effects from Unconventional Natural Gas Development. *Env. Sci. Tech*, in press.
- (25) Roy A. A., Adams, P. J., Robinson A. L., (2014) Air pollutant emissions from the development, production, and processing of Marcellus Shale natural gas. *Journal of the Air & Waste Management Association*, 64, 19-37.
- (26) Rail Road Commission of Texas, *Texas Gas Well Gas Production in the Newark, East (Barnett Shale) Field – 1993-2011*. Retrieved 9/11/2012 from http://www.rrc.state.tx.us/barnettshale/NewarkEastField_1993-2011.pdf

(27) Gilman, J. B., Lerner B. M., Kuster, W.C., de Gouw, J. A. (2013) Source Signature of Volatile Organic Compounds from Oil and Natural Gas Operations in the Northeastern Colorado. *Environmental Science and Technology*, 47, 1297-1305.

(28) Edwards, P.M., Young, C. J., Aikin, K., deGow, J., Dube, W. P., Geiger, F., Gilman, J., Helmig D., Holloway, J. S., Kercher, J., Lerner, B., Martin, R., McLaren R., Parrish, D. D., Peischl J., Roberts, J.M., Ryerson T. B., Thornton, J., Warneke C., Williams, E. J., Brown, S., S., (2013) Ozone photochemistry in an oil and natural gas extraction region during winter: simulations of a snow-free season in the Uintah Basin, Utah, *Atmos. Chem. Phys.*, 13, 9955-8971.

3 METHANE EMISSIONS FROM PNEUMATIC CONTROLLERS

3.1 Introduction

As mentioned in the previous chapter, Methane emissions in the natural gas supply chain have been estimated using two basic approaches, commonly referred to as top-down and bottom-up approaches. Top-down approaches for estimating methane emissions from the natural gas supply chain involve measuring ambient concentrations of methane near emission sources. These concentrations can be measured using fixed ground monitors [1, 2], mobile and vehicle mounted ground monitors [3, 4], aircraft based instruments [5-7] or satellite instruments [8]. Brandt *et al.* [9] and Miller, *et al.* [10] have summarized recent top-down estimates of methane emissions and conclude that top-down emission estimates are generally higher than current bottom-up inventories of methane emissions, and some of this difference may be due to methane emissions from the natural gas supply chain. However, these analyses do not reveal which of the many potential sources of methane emissions along the natural gas supply chain might be incorrectly estimated.

Complementing top-down measurements, bottom-up measurements of methane emissions are made directly at the emission sources. In this approach, emission measurements are made at a representative sample of sources; the measurements from the sample population are then extrapolated to larger regional or national populations. The advantage of “bottom-up” approaches is that they can gather much more detail about the emission sources, and therefore can identify which source categories, among many, are responsible for emissions. For example, Allen, *et al* [11] concluded that emissions from well completion flowbacks are over-estimated, while emissions from pneumatic controllers may be under-estimated, in current inventories of emissions. Both top-down and bottom-up approaches can contribute to an improved understanding of methane emissions from the natural gas supply chain.

The work reported in the present chapter uses bottom-up measurements to improve understanding of emissions from pneumatic controllers on natural gas production sites.

Similarly, *Chapter 4* uses bottom-up measurements to improve understanding of emissions from liquid unloadings of natural gas wells.

Pneumatic controllers use gas pressure to control the operation of mechanical devices, such as valves. The valves, in turn, control process conditions such as levels, temperatures and pressures. When a pneumatic controller identifies the need to change liquid level, pressure, temperature or flow, it will open or close a control valve in order to return to a desired set point. The opening and closing of the valve can occur either through discrete (on/off) changes, or through changes that are proportional in magnitude to the deviation from the set point (throttling). Controllers can deliver this type of service (on/off and throttling) through either continuously venting or intermittent venting of gas. Thus, controllers can be grouped into four categories, depending on the type of service (on/off or throttling) and the type of venting (continuous or intermittent). In estimating emissions, the U.S. EPA uses the categories of low continuous bleed (<6 scf/h of gas vented), high continuous bleed (>6 standard cubic feet per hour (scf/h) of gas vented) and intermittent controllers [12]. Finally, controllers can also be categorized based on equipment manufacturer, model number, and the type of application (e.g., separator level control) in which they are used. In this work, the primary categorization of controllers will be as either continuous vent or intermittent vent; data on applications, service types, and EPA categorization for the controllers sampled in this work are provided in the Appendices (Section A.4).

The U.S. EPA [12] reports 447,600 pneumatic controllers are in use at natural gas production sites in the United States. These controllers are estimated to emit 334 Gg/yr of methane (17.4 billion cubic feet (bcf) methane), for an average of 0.7 Mg methane device⁻¹ yr⁻¹ or 4.2 scf/h methane device⁻¹. These estimated emissions from pneumatic controllers have been based on relatively limited measurements [13]; recent field measurements have suggested that these emissions may be understated.

Allen *et al.* [11] made measurements of emissions from 305 pneumatic controllers on well sites in the United States where the wells had been hydraulically fractured. Average emissions were 10.5 scf/h of methane, approximately double the average emission rate per device in the current EPA national emission inventory. Measurements of emissions from 581 pneumatic controllers, made in British Columbia and Alberta, averaged 9.2 scf/h of whole gas [14], an emission rate similar to that reported by Allen, *et al.* [11]. In both of these studies, emissions from controllers exhibited wide ranges. In both sets of measurements, a small subset of controllers accounted for most of the emissions.

While the measurements at hydraulically fractured gas wells in the United States [11] and the measurements in British Columbia and Alberta [14] recorded emissions higher than the average emissions per device in the EPA national emission inventory, the sample sets for these two sets of measurements were not necessarily broadly representative of U.S. national populations of pneumatic controllers. The measurements reported by Allen, *et al.* [11] were made exclusively in shale gas production regions, and at sites where the wells had been hydraulically fractured. Many of the sites were recently completed wells, which initially tend to have higher liquid production rates, and therefore may have more frequent actuation of certain types of pneumatic devices than the average for the entire population of gas wells in the United States, leading to potentially higher emissions. The Canadian measurements [14] were made exclusively in one production region and on devices with manufacturer specified emission rates in excess of 4.2 scf/h.

The goals of the work presented in this chapter were (i) to measure emissions from pneumatic controllers at a wider population of wells, geographically distributed across the United States, including conventional gas wells, shale gas wells and a limited number of oil wells, and (ii) to characterize the features of the controllers with high emissions, which previous work [11, 14] has found to be the major contributor to the emissions.

3.2 Methods

3.2.1 Sampled Population

A total of 377 pneumatic controllers were sampled at 65 sites (some with multiple wells) throughout the United States (an average of 5.8 pneumatic controllers per site, 2.7 controllers per well). Measurements were made primarily at natural gas production sites (351 of 377 controllers), and at a limited number of oil sites (26 controllers). Because the definitions of oil and gas wells vary, largely depending on gas to oil production ratios, the data will be treated as a single set. Sampling sites were selected randomly from well sites owned by companies participating in the study (see Appendices, Section A.1). For each well site that was visited, all controllers on the site were sampled, unless operating conditions or safety issues prevented sampling. The applications that the controllers were used in (e.g., separator level control, compressor, pressure control) are shown in Table 3-1. Details of the regions, device types, associated well types, operating methods and other characteristics of each of the 377 controllers sampled in this work are provided in Appendices, Section A.4.

Table 3-1. Sample population, categorized by region and controller application

Region	Application								Total
	Sep.	Proc. Heater	Comp.	Wellhead	Plunger Lift	Dehyd. Syst.	Flare	Sales	
AP	14	13	0	24	1	0	0	0	52
GC	73	0	13	11	7	16	1	1	122
MC	48	11	7	0	11	1	0	0	78
RM	51	21	0	32	11	8	2	0	125
Total	186	45	20	67	30	25	3	1	377

3.2.2 Emission measurement methods

Emissions from pneumatic controllers can be determined either by measuring the supply of gas entering the controller or by measuring the gas discharged from the controller. Both approaches were used in this work, and since there is no accumulation of gas in the

controller, both measurement approaches should lead, in the absence of emissions from the control valve, to equivalent measurements.

Measurements of the gas entering the controller were made by one of three Fox flow meters (Model #FT2A); flow meters were inserted into the supply gas line for the controller. This supply gas measurement was the primary measurement method used in this work, and was used to measure emission rates on 333 of the 377 controllers in the sample population (the remainder were sampled by measuring gas emitted by the controller, see Appendices, section A.3). The flow meters reported flows at a sampling frequency of 10 Hz. Two of the Fox Model #FT2A instruments (labeled A and C in this study) had a range of operation of 0-300 scf/h, with a precision of $\pm 1\%$ of flow, and the third Fox Model #FT2A (labeled B in this study) had a range of operation of 0-1200 scf/h, with a precision of $\pm 1\%$ of flow. The Fox model #FT2A instruments A and C were used whenever possible because of their greater absolute precision, however, if any instantaneous reading on the A or C Fox Model #FT2A was greater than 300 scf/h, the measurement on the pneumatic device was repeated with the B meter to ensure that high leak rates were measured accurately. This happened only once during the measurement campaign, and for this single controller, the flow exceeded 300 scf/h only during a few seconds when the flow from an actuation was peaking (average whole gas flow rate over fifteen minutes of sampling was 3.06 scf/h). A repeat test with the B instrument did not detect any actuations.

For each controller measurement using the supply gas flow meter, a site operator depressurized and disconnected the controller supply gas line; the flow meter was inserted and the system was reconnected, re-pressurized, and allowed to stabilize for several minutes before measurements began. Once the system had stabilized, measurements were made for approximately 15 minutes. Longer sampling times may have allowed a more complete measurement of emission rates from devices with relatively fewer controller actuations, but would have limited the number of controllers

that could be sampled. Figure 3-2 shows representative 15 minute emission time series for pneumatic controllers measured using the supply gas measurement.

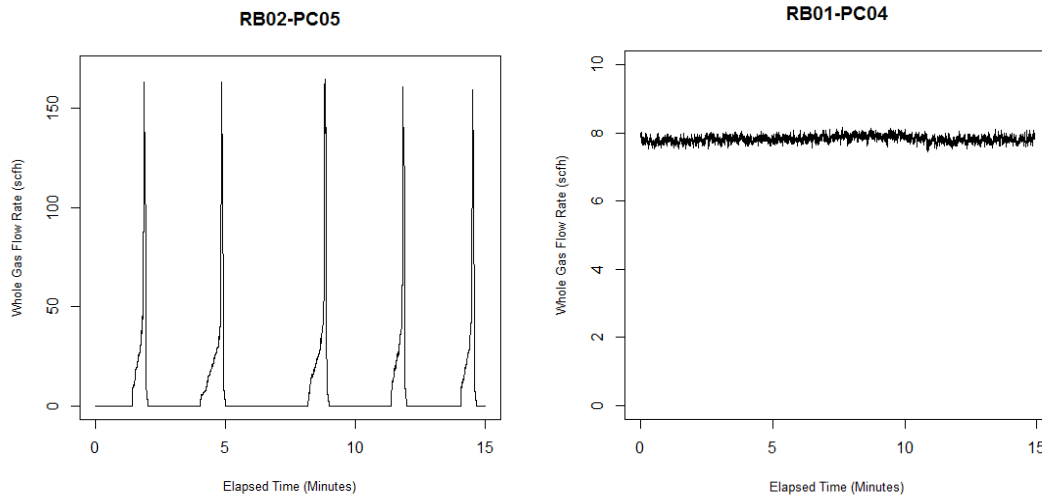


Figure 3-1. Representative time series for supply gas measurements for intermittent vent and continuous vent controllers; the actuating controller (RB02-PC05) had a total of 5 actuations during the sampling period and an average emission rate, over the 15 minute period of 7.9 scf/h; the throttling controller (RB01-PC04) had nearly constant emissions of 8.0 scf/h.

All three Fox flow meters were calibrated by the instrument manufacturer and tested in the laboratory, using methane. The instruments measure flow based on a thermal conductivity measurement. In this work, since gas composition information was available for each site where measurements were made, site specific correction factors were employed to estimate methane and whole gas emission rates. The method is described in the Appendices (see Section A.2). Results in this work are reported as both methane and whole gas emission rates, based on site specific gas composition data.

For some pneumatic controllers, it was not possible or safe to disrupt the supply gas to insert the supply gas flow meter, so exhaust gas measurements were used as the primary measurement on that subset of devices. Exhaust gas flow rate was measured using a Hi Flow® instrument similar to that described by Allen *et al.* [11]. Briefly, the Hi Flow® Sampler is a portable, intrinsically safe, battery-powered instrument that has been used

for several decades in measuring emissions of methane in the natural gas supply chain [15-17]. An emission source is enclosed, using attachments that come with the instrument; leak rate is measured by drawing air from the enclosure, through the sampler, at a high flow rate (up to 8-10 cfm) to capture all the gas emitted by the component, along with a certain amount of entrained surrounding air. By accurately measuring the flow rate of the sampled stream and the background corrected natural gas concentration within the sampled stream, the gas leak rate is calculated. Methane is measured, at concentrations less than approximately 5%, by a catalytic oxidizer unit coupled with a thermal conductivity detector. At methane concentrations greater than approximately 5%, concentrations are measured directly using a thermal conductivity detector. The instrument was calibrated in the field using pure methane and a mixture containing 2.5% methane. The instrument reading based on the methane calibration was corrected for gas composition using site specific gas composition data and laboratory data, as described in Appendices (A.2). The commercial Hi Flow® instrument is designed primarily to measure methane leaks that have a relatively steady flow and flow rates are not normally automatically recorded at high frequency. For this work, the instrument software was modified by the manufacturer to output data every 2-3 seconds. A 0.3-0.5 Hz reporting frequency was selected based on residence times expected in the leak enclosures at the maximum flow rate of the Hi Flow® device (at a 10 cubic feet per minute sample flow, gas in a 1 ft³ sample enclosure has a residence time of 6 seconds). As with the in-line supply gas measurement, Hi Flow® data were collected for approximately 15 minutes for each controller.

A time series from the Hi Flow® device, along with a parallel measurement made using a supply gas meter, is shown in Figure 3-3. The Hi Flow® device, because it entrains ambient air in a long sample loop, dampens some of the peak rate. Therefore, the Hi Flow® is not able to resolve high frequency actuations as well as the in-line supply measurement. For 24 controller measurements, both supply gas and Hi Flow® measurements were made. The detailed results are provided in Appendices (Section

A.3). To summarize, 11 of the 24 simultaneous measurements had emissions of less than 0.005 scfh (46%). For 5 devices which had an average emission rate greater than 6 scfh (measured by the supply gas meter) the supply gas meter to Hi Flow® measurement ratio was between 0.7 and 1.1.

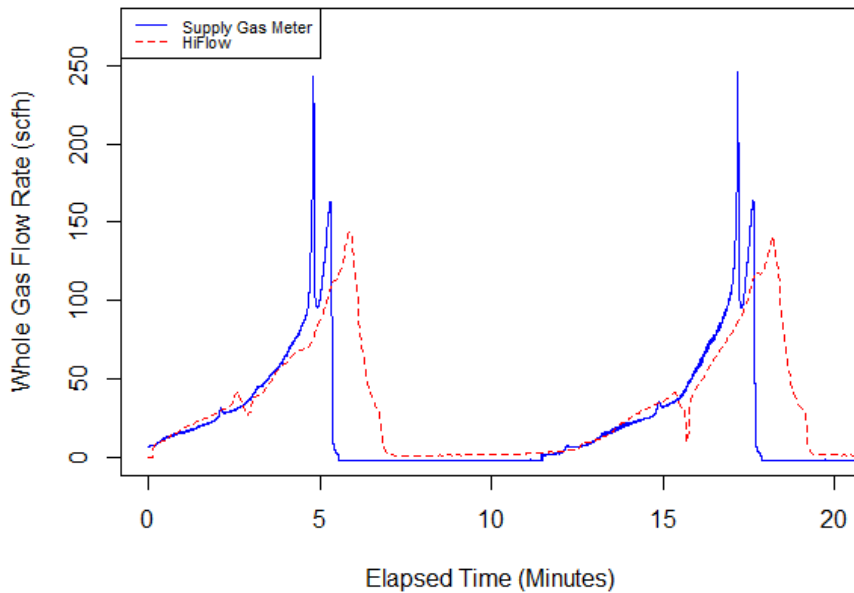


Figure 3-2. Comparison of supply gas meter (blue line) and Hi Flow® measurements (red line) for device LB07-PC04, which was a water level control on a separator. The average emission rate measured by the supply gas meter was 27.0 scf/h as compared to 33.9 scf/h measured by the Hi Flow®. Note that the time lag, longer period of emission detection, and the reduced maximum flow rate associated with the Hi Flow® measurement is expected because of the dilution that occurs with ambient air in the exhaust enclosure and the flow through the instrument.

3.3 Results and Discussion

Methane emissions from 377 controllers were measured in this work and details of each of the individual measurements are available in Appendices (Section A.4). A relatively small subset of devices accounts for a majority of the emissions. At the high end of the emission rate distribution, twenty percent of devices accounted for 96% of whole gas and methane emissions. The 19% of devices that had emissions in excess of 6 standard cubic

feet whole gas per hour (scf/h) accounted for 95% of all whole gas and methane emissions. At the low emission rate end of the distribution, more than half (51%) of the controllers had an emissions rate less than 0.001 scf/h over the 15 minute sampling period; 62% had an emissions rate less than 0.01 scf/h over the 15 minute sampling period.

The average emission rate for the 377 devices is 5.5 scf/h of whole gas (4.9 scf/h of methane), however, this average emission may be influenced by the estimated emission rates for devices that had no emissions over the 15 minute sampling period. If the devices with no emissions detected over 15 minutes are assigned the lowest emission rate detected (0.001-0.01 scf/h), there is no change in the average emission rate. However, using this minimum detection limit approach may under-estimate potential emissions for devices that had little to no detectable emissions over 15 minutes. Some of these devices may have actuations with relatively large volumes, but that are relatively infrequent. For example, the device that generated the data shown in Figure 3-3 had emissions of more than 2 cubic feet of gas associated with a single actuation, and in principle, a device could actuate up to four times per hour and not be detected over a 15 minute sampling period. Therefore, to estimate the emissions from devices with no emissions detected over a 15 minute sampling period, the average emission per actuation was calculated for controllers in each application. The average emissions per actuation were multiplied by an estimated frequency of actuation. For example, for separator level controllers, the average volume per actuation was estimated by averaging observed volumes per actuation for separator level controllers; the average frequency of actuation for devices, for which no actuations were observed, was estimated by extrapolating observed actuation frequency data for controllers in separator level control service. Details are available in Appendices (Section A.5). Using this approach, the estimated average emissions associated with devices with no emissions recorded over a 15 minute sampling increases the population average emissions by 2-6%. Because this increase is relatively small, for clarity, all of

the data reported in this work are based on actual measurements, not including additions to the emissions for devices with low (0.001-0.01 scf/h) observed emissions.

To estimate an uncertainty bound on the overall average, a bootstrapping process was used [18]. In the bootstrapping procedure, the original data set of 377 devices was recreated by making 377 random device selections, with replacement, from the data set. A total of 1000 of these re-sampled data sets were created and the mean value of the emissions for each re-sampled data set was determined. The 95% confidence interval for the whole gas emission estimate of 5.5 scf/h is 4.0-7.2 scf/h, where the bounds represent the 2.5% and 97.5% percentiles of the means in the 1000 re-sampled datasets. Similarly, the 95% confidence interval for the methane emissions estimate of 4.9 scf/h is 3.6-6.5 scf/h.

The measurements showed significant variations among regions, the controller application, and whether the device was continuous vent or intermittent vent. Table 3-2 summarizes the distribution of emission rates among controllers in various applications, and shows the regional distribution of controller emissions. Measurements made on pneumatics in service on compressors had average emission rates of 14.0 scf/h (12.4 scf/h methane), compared to an average whole gas emission rate of 5.5 scf/h (4.9 scf/h methane) for all devices. Devices in use for level control on separators averaged 8.1 scf whole gas/h (7.1 scf methane/h). Overall, 76% of devices measured with whole gas emission rates greater than 6 scf/h were in service on compressors or as level controllers on separators. Emission rates for continuous vent controllers (57 devices, average emissions of 24.1 scf/h whole gas, 21.8 scf/h methane) were higher than for intermittent devices (2.2 scf/h whole gas, 1.9 scf/h methane).

Table 3-2. Whole gas emissions from controllers (scf/h), categorized by region and application.*

Average whole gas emission rates from controllers (scf/h), categorized by the application

Region	Average whole gas emission rates from controllers (scf/h), categorized by the application									
	All Devices	Separator	Process Heater	Compressor	Wellhead	Plunger Lift	Dehydration System	Flare	Sales	Avg. w/o Compressors
AP	1.7	0.3	1.3	-	2.8	0.0	-	-	-	1.7
GC	11.9	16.3	-	10.6	0.0	7.3	4.4	0.0	0.0	12.0
MC	5.8	4.9	0.0	20.2	-	6.5	4.2	-	-	4.4
RM	0.8	1.5	0.2	-	0.4	0.1	0.0	0.0	-	0.8
Total	5.5	8.1	0.5	14.0	1.2	4.1	3.0	0.0	0.0	5.0

*Numbers of devices sampled in each category are reported in Table 3-1.

In addition to varying by application and controller type (continuous vent or intermittent vent), emissions also varied by region. Emissions were highest in the Gulf Coast and Mid-Continent regions and were lowest in the Rocky Mountain and Appalachian regions (see Appendices, Section A.4, for geographical boundaries of regions). Controllers on compressors, with high average emissions, were only observed on sampling sites in the Gulf Coast and Mid-Continent regions, so some of the regional differences can be attributed to the presence of compressors. As shown in Table 3-2, however, if average emissions by region are recalculated without including controllers associated with compressors, the Gulf Coast and Mid-Continent regions still had average emissions greater than those observed in the Rocky Mountain and Appalachian regions.

Another factor that may account for regional differences in emission rates is frequency of actuation. For example, controllers on separators in the Gulf Coast could actuate more frequently due to higher liquid production rates, which could explain higher emission rates in that region. However, the frequencies of actuation for the devices in Gulf Coast were similar to those in most other regions, indicating a larger emission per actuation for the devices in the Gulf Coast, rather than more frequent actuation. In contrast, the Appalachians showed a considerably higher frequency of actuations and a smaller

emission rate, indicating a smaller emission per actuation for those devices (Table 3-3). Thus, regional differences in pneumatic controller emission rates cannot be completely explained by frequency of actuation of controllers, or by controllers associated with compressors and separator level control (Table 3-2); much of the difference may be due to differences in controller type (continuous vent vs. intermittent vent) among regions. Continuous vent devices, with average whole gas emissions of 24.1 scf/h, were 21% of the controllers in the Gulf Coast and Mid-Continent regions, but only 9% in the Appalachian and Rocky Mountain regions.

Table 3-3. Frequency of actuations and emissions from on-off controllers where actuations were observed, categorized by region

Region	Count of Devices	Frequency of Actuation (#/min)	Avg. Whole Gas Emission rate (scf/h)
AP	8	2.42	4.85
GC	29	0.39	21.02
MC	18	0.94	4.97
RM	25	0.43	1.72
Total	80	Average: 0.73	Average: 9.76

This data set of emissions from pneumatic devices has elements that are similar to and different from the previous data sets reported for the United States [11], and for British Columbia and Alberta [14]. The primary similarity is that all three data sets indicate that a small population of devices dominates total emissions. In this work, 19% of devices with emissions greater than 6 scf/h of whole gas account for 95% of the whole gas and methane emissions. In the previous measurements reported by Allen, *et al.* [11], 20% of devices account for 80% of the whole gas emissions and 41% of devices with emissions greater than 6 scf/h of whole gas, account for 90% of the whole gas emissions (88% of the methane emissions). In the measurements for British Columbia and Alberta [14] (referred to here as the British Columbia data), which were restricted to pneumatic devices with manufacturer reported bleed rates greater than 4.2 scf/h, 44% of devices with emissions greater than 6 scf/h accounted for 91% of emissions. Both the British Columbia data and the measurements reported in this work had large numbers of devices for which no emissions were detected during the sampling period. For the British Columbia data (again, focused on devices with manufacturer reported bleed rates in excess of 4.2 cfh), 31% of measurements had no detectable emissions over a 30 minute sampling period; in this work 62% of devices had emissions less than 0.01 scf/h over the 15 minute sampling period.

The overall average emission rates reported in this work are lower than the previous data sets reported for the United States [11], and for British Columbia and Alberta [14]. For

the British Columbia data this can be attributed to the sampling design for that data set, which selected devices with manufacturer reported bleed rates in excess of 4.2 scf/h. These controller types tend to be found in particular applications. When the emissions from the British Columbia data set are compared to the emissions reported in this work, for devices in similar applications, the results are in reasonable agreement. For example, for the separator controllers that were the most frequent application observed in this work, the British Columbia data report an average emissions rate of approximately 7.8 scf/h (level control) while the average for this work was 8.1 scf/h (separator application).

The lower average emission rates reported in this work, compared to those reported by Allen, *et al.* [11] is primarily due to the number of controllers with no emissions detected over the sampling period. Figure 3-4 compares emission rates for controllers sampled in this work, with emissions rates reported by Allen, *et al.* [11]. The results show reasonable agreement between the two studies for controllers with emissions above 6 scf/h. These controllers accounted for 95% of the emissions in this work and 90% of the emissions in the sample reported by Allen, *et al.* [21].

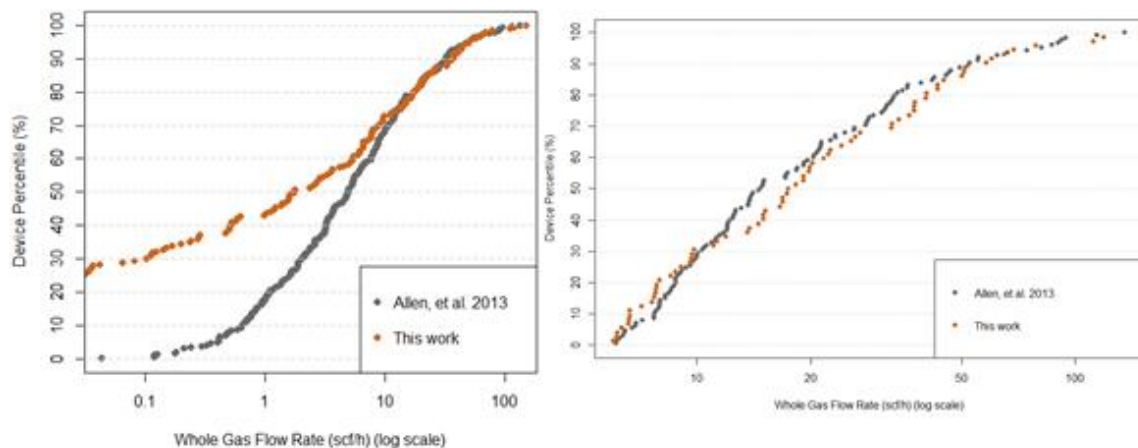


Figure 3-3. Distributions of emissions for subsets of controllers venting greater than 0.01 (left) and subsets of controllers venting greater than 6 scf/h of whole gas as reported in this work and Allen, *et al.* [11].

The primary reason for the differences in the average emission rates reported in this work and in Allen, *et al.* [11] is the higher percentage of low emission devices (<0.01 scf/h) observed in this work. This could be due to multiple factors. In this work, all controllers on-site were sampled, regardless of whether they would be reported through emission inventories. For example, Emergency Shut-Down (ESD) controllers represented 12% of the sampled population in this work. These controllers do not have planned actuations, so they would not have been sampled in the work of Allen, *et al.* [11], and they may or may not be included in controller counts in national emission inventories. In addition, in the work of Allen, *et al.* [11] about 40% of the inventoried controllers on sites were sampled; while these were intended to be selected randomly from inventoried controllers, there may have been an unintentional bias toward devices that were observed, with an infrared camera, to have emissions.

3.4 Implications for national emission estimates

As shown in Table 3-4, if regional average emission rates determined in this work are multiplied by regional controller counts reported in the 2012 EPA national greenhouse gas emission inventory (2012 GHG NEI, released in 2014 [12]), the national methane emission estimate for pneumatic controllers in natural gas service is 313 Gg/yr (within 10% of the 2012 GHG NEI estimate of 334 Gg). If the national average of the emission rates measured in this work (5.5 scf/h of whole gas, or 4.9 scf/h of methane) is multiplied by the total national equipment count (447,606 controllers) the national methane emission estimate is 369 Gg/yr, 10% higher than the 2012 GHG NEI estimate of 334 Gg [12]. Adding an additional 2-6% to the estimated emission totals to account for potential emissions from controllers that had less than 0.01 scf/hr of emissions over 15 minutes, would only slightly change these comparisons with the 2012 GHG NEI. This estimate may represent a lower bound on national emissions, however, since the average emissions per controller observed in this work includes some low emitting devices, such as ESD controllers, that may not be included in the count of national controllers. If the

average emissions per controller from this work were recalculated with ESD controllers excluded, the average emissions would increase by approximately 15% (see Appendices, Section A.7).

The inclusion or exclusion of ESD controllers in national pneumatic controller counts is just one part of the uncertainty associated with the total count of controllers. The average number of controllers per well observed in this work (2.7 controllers per well) was higher than the average number of controllers per well (1.0 controllers per well) reported in the 2012 GHG NEI, potentially indicating an under-count of controllers in the GHG NEI. Some of the difference between the controllers per well observed in this work and the average pneumatic controllers per well in the GHG NEI is due to wells that use mechanical or other non-pneumatic controllers, as an example of how an alternative controller count could influence national emission estimates, if 75% of wells in the United States have an average of 2.7 pneumatic controllers per well (the remainder having non-pneumatic controllers), the total count of pneumatic controllers would double the level in the current inventory, roughly doubling emissions. It was beyond the scope of this work to develop new national pneumatic controller counts, but the data reported here indicate that this is a topic that merits attention.

Table 3-4. National emission estimates, based on regional device counts for pneumatic controllers and regional average emissions measured in this work.

Region	Count of Devices	Avg. Emission rate whole gas(scfh)	Avg. Emission rate methane (scfh)	Regional emissions (Gg/yr)
AP	77,261	1.70	1.65	21.5
GC*	53,436	11.86	10.67	95.9
MC	222,684	5.78	4.85	181.8
RM**	124,225	0.75	0.67	14.0
Total	447,606			313.2

*MC totals include equipment counts for Mid-Continent and Southwest regions reported in the 2012 EPA GHG NEI

** RM totals include equipment counts for Rocky Mountain and West Coast regions reported in the 2012 EPA GHG NEI

3.5 Characteristics of high emitting devices

Because average emissions are strongly influenced by the highest emitting devices, the characteristics of the 40 controllers with highest emissions rates were examined in detail by experts in pneumatic device operation. These characterizations included the service type, region of use, device type, number of actuations, and other temporal features of the emission time series. Based on these analyses, many of the devices in the high emitting group were behaving in a manner inconsistent with the manufacturer's design. For example, some devices not designed to bleed continuously had continuous emissions. This could be the result of a defect in the system, such as a crack or hole in the end-device's (control valve's) diaphragm actuator, or a defect in the controller itself, such as fouling or wear. No additional troubleshooting analysis was performed on these high emitters, so the actual root causes are not known with certainty. The results, however, do indicate that some of the high emissions were caused by repairable issues. Details are provided in the Appendices (A.8).

3.6 Conclusions

Emissions from 377 gas actuated (pneumatic) controllers were measured at natural gas production sites and a small number of oil production sites, throughout the United States. A small subset of the devices (19%), with whole gas emission rates in excess of 6 standard cubic feet per hour (scf/h), accounted for 95% of emissions. More than half of the controllers recorded emissions of 0.001 scf/h or less during 15 minutes of measurement. Pneumatic controllers in level control applications on separators and in compressor applications had higher emission rates than controllers in other types of applications. Regional differences in emissions were observed, with the lowest emissions measured in the Rocky Mountains and the highest emissions in the Gulf Coast. Average emissions per controller reported in this work are 17% higher than the average emissions per controller in the 2012 EPA greenhouse gas national emission inventory (2012 GHG

NEI, released in 2014); the average of 2.7 controllers per well observed in this work is higher than the 1.0 controllers per well reported in the 2012 GHG NEI.

3.7 References

(1) Zavala-Araiza D, Sullivan DW, and Allen DT (2014). Atmospheric hydrocarbon emissions and concentrations in the Barnett Shale natural gas production region. *Environ. Sci. Tech.*, doi: 10.1021/es405770h.

(2) Katzenstein AS, Doezema LA, Simpson IJ, Blake DR, Rowland FS (2003) Extensive regional atmospheric hydrocarbon pollution in the southwestern United States. *Proc. Natl. Acad. Sci. U.S.A.* 100: 11975–11979.

(3) Herndon SC, Floerchinger C, Roscioli JR, Yacovitch TI, Franklin JF, Shorter JH, Kolb, CE, Subramanian R, Robinson AL, Molina LT, *et al.* (2013) Measuring methane emissions from industrial and waste processing sites using the dual tracer flux ratio method Annual Meeting the American Geophysical Union, San Francisco, December 2013.

(4) Herndon SC, Jayne JT, Zahniser MS, Worsnop DR, Knighton B, Alwine E, Lamb BK, Zavala M, Nelson DD, McManus JB, *et al.* (2005) Characterization of urban pollutant emission fluxes and ambient concentration distributions using a mobile laboratory with rapid response instrumentation *Faraday Discuss.* 130: 327–339.

(5) Petron G, Frost G, Miller BR, Hirsch AI, Montzka SA, Karion A, Trainer M, Sweeney C, Andrews AE, Miller L, *et al.* (2012). Hydrocarbon Emissions Characterization in the Colorado Front Range: A Pilot Study *J. Geophys. Res.* 2012, 117(D4): D04304.

(6) Peischl J, Ryerson TB, Brioude J, Aikin KC, Andrews AE, Atlas E, Blake D, Daube BC, de Gouw JA, Dlugokencky E, *et al.* (2013) Quantifying Sources of Methane Using

Light Alkanes in the Los Angeles Basin, California *Journal of Geophysical Research Atmospheres* 2013, 118:4974-4990.

(7) Karion A, Sweeney C, Pétron G, Frost G, Hardesty RM, Kofler J, Miller BR, Newberger T, Wolter S, Banta R, *et al.* (2013) Methane emissions estimate from airborne measurements over a western United States natural gas field *Geophysical Research Letters* 2013, 40: 1–5, doi:10.1002/grl.50811.

(8) Wecht KJ, Jacob DJ, Sulpizio MP, Santoni GW, Wofsy SC, Parker R, Bosch H. and Worden J. (2014). Spatially resolving methane emissions in California: constraints from the CalNex aircraft campaign and from present (GOSAT, TES) and future (TROPOMI, geostationary) satellite observations. *APCD*, 14: 4119-4148.

(9) Brandt AR, Heath GA, Kort EA, O’Sullivan F, Pétron G, Jordaan, SM, Tans P, Wilcox J, Gopstein, AM, Arent D, *et al.* (2014). Methane Leaks from North American Natural Gas Systems. *Science*, 343: 733-735.

(10) Miller SM, Wofsy SC, Michalak AM, Kort EA, Andrews AE, Biraud SC, Dlugokencky EJ, Eluszkiewicz J, Fischer ML, Janssens-Maenhout G, *et al.* (2013) Anthropogenic emissions of methane in the United States. *Proc. Natl. Acad. Sci. U.S.A.*, 110, 20018–20022.

(11) Allen DT, Torres VM, Thomas J, Sullivan D, Harrison M, Hendler, A., Herndon, SC, Kolb, CE, Fraser, MP, Hill, AD, *et al.* (2013) Measurements of Methane Emissions at Natural Gas Production Sites in the United States, *Proc. Natl. Acad. Sci. U.S.A.*, 110: 17768-17773.

(12) U.S. Environmental Protection Agency (EPA). Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2012. April 2014.

(13) U.S. Environmental Protection Agency (EPA), Office of Air Quality Planning and Standards. Oil and Natural Gas Sector Pneumatic Devices; Report for Review Panel, April 2014.

- (14) Prasino Group, Final Report for determining bleed rates for pneumatic devices in British Columbia. Report to British Columbia Ministry of Environment, December 2013.
- (15) Harrison, M.R.; Shires, T.M.; Wessels, J.K.; Cowgill, R. M; Methane Emissions from the Natural Gas Industry, Volumes 1 – 15, Final Report, GRI-94/0257 and EPA-600/R-96-080, Gas Research Institute and US Environmental Protection Agency, June 1996.
- (16) Harrison, M.R.; Shires, T.M.; Wessels, J.K.; Cowgill, R.M.; Methane Emissions from the Natural Gas Industry, EPA/600/SR-96/080 June 1997.
- (17) Kirchgessner, D.A.; Lott, R.A. Cowgill, R.M.; Harrison, M.R.; Shires, T.M. Estimate of Methane Emissions from the U.S. Natural Gas Industry, Chemosphere, 35, 1365-1390, 1997.
- (18) Efron B.; Tibshirani, R. An Introduction to the Bootstrap, Chapman & Hall, CRC Press, (1993).

4 METHANE EMISSIONS FROM LIQUIDS UNLOADINGS

4.1 Introduction

Measurements of emissions from pneumatic controllers have been described in the previous chapter. This work reports on emissions from gas well liquid unloadings.

A liquid unloading may be necessary when a gas well that also produces oil or water accumulates liquids in the well bore. The liquids accumulation may be due to a variety of causes, including decreases in gas velocity in the well, decreases in reservoir pressure, or changing gas to liquid ratios. As liquids accumulate, well production can decline and an operator may choose to unload the liquids from the well to restore production. Liquids can be unloaded in a variety of ways. For example, the well tubing can be modified to increase gas velocity or a pump may be installed to remove downhole liquids. Neither of these methods lead to venting emissions. Other unloading methods, such as temporarily diverting the flow from the well to an atmospheric vent, do lead to emissions. This work focuses on unloadings that result in emissions.

In the most recent national inventory of greenhouse gas emissions (for calendar year 2012, released in 2014, referred to here as the EPA 2012 GHG NEI) [1], the EPA estimates that 60,810 natural gas wells, out of an estimated 470,913 natural gas wells in the United States (not including oil wells with associated gas production), have liquid unloadings that result in methane emissions. This represents 13% of gas wells in the EPA 2012 GHG NEI. Collectively, liquid unloadings from these wells are estimated to emit 273.6 Gg of methane per year (14.2 billion standard cubic feet, bcf), or approximately 14% of the estimated 1,992 Gg of methane emissions from the natural gas production portion of the natural gas supply chain.

The estimates of methane emissions from liquid unloadings in EPA 2012 GHG NEI are generally consistent with more recent information collected through the EPA's Greenhouse Gas Reporting Program (for calendar year 2012, released in 2013, referred to here as the EPA 2012 GHGRP) [2]. The GHGRP reports approximately 276 Gg of

methane emissions from liquid unloadings at facilities that meet threshold reporting requirements. Information for 58,663 wells that have unloading emissions was reported in 2012. The liquid unloading emission estimates from the EPA GHGRP can be disaggregated by production region and these data are shown in Appendices (*Section B.1*).

Emissions from liquid unloadings of natural gas wells are not uniformly distributed in time or space. Estimated emissions from liquid unloadings are spatially concentrated in Rocky Mountain production regions. Wells in the western United States account for more than half of estimated emissions from liquid unloadings in the 2012 GHGRP. Temporal distributions also vary. Some wells release unloading emissions several times per day while others may release unloading emissions only once per year or once during the well's production life cycle. Wells may only release unloading emissions for a portion of their production lifetime, leading to a dependence of unloading emissions on well age. In addition to spatial and temporal variability in emissions of wells that vent, both estimates and measurements indicate that a small fraction of wells that vent account for a majority of emissions. For example, for one type of well with unloading emissions (wells without plunger lifts – see definition later in text), emission estimates reported by the American Petroleum Institute/America's Natural Gas Alliance (API/ANGA), indicate that three percent of wells accounted for half of emissions from this type of well and half of the wells accounted for more than 90% of emissions [3]. In a limited number of measurements of methane emissions from a single type of well with unloading emissions (wells without plunger lifts – see definition later in text), Allen, *et al.* [4] found that 95% of the emissions came from less than half of the wells.

Emission estimates, and a limited number of measurements of methane emissions from liquid unloadings, both suggest that a small fraction of wells, in particular geographical regions, and at particular times in the well's life cycle, account for a large fraction of liquid unloading emissions. These characterizations of unloading emissions are primarily based on emission estimates, however, and there are few data in the scientific

literature to test the reliability of emission estimates. This leads to potentially large uncertainties in the emissions from this source category. More measurement data are needed, along with a better understanding of the relationships between well characteristics and unloading emissions.

This work reports measurements of methane emissions from 107 natural gas wells with emissions associated with liquid unloading. These data represent the most extensive set of measurements of emissions from liquid unloadings in the scientific literature. The relationships between emissions magnitude, unloading event frequency and other well characteristics are explored.

4.2 Methods

4.2.1 Emission Measurements

The liquid unloadings reported in this work are grouped as plunger-lift unloadings and unloadings of wells without plunger lifts.

In a manually triggered unloading of a well without a plunger lift, an operator manually diverts the well's flow from a production separator, which typically operates at pressures of multiple atmospheres, to an atmospheric pressure tank. This allows the well to flow to a lower pressure destination (the atmospheric pressure tank or vent, rather than the pressurized separator). The resulting higher pressure gradient allows more gas to flow, increasing velocity in the production tubing and entraining and lifting liquids out of the well. Gas is discharged through the tank vent to the atmosphere. In a small number of wells (~0.1% of wells reported by companies participating in this work), this process is automated, resulting in two sub-categories of unloadings for wells without plunger lifts, manual and automatic. All of the measurements reported in this work for wells without plunger lifts are for wells that had unloadings that were manually triggered; no wells without plunger lifts were observed in the sampling that had automated unloadings.

Emissions from unloadings of wells without plunger lifts were measured in this work by directing flow through a temporary stack installed on top of the vent. Figure 4-1 shows a conceptual diagram of a tank layout on a well site and the positioning of the temporary stack. Grounded metal or metal lined tubing was used to construct the temporary stack, to prevent static discharge. Flow rate through the temporary stack was measured continuously, near the centerline of the temporary stack, using a thermal gas mass flow meter. The thermal meter was extended into the middle of the temporary stack, which was between two and eight inches in diameter, with the diameter depending on the anticipated flow rate. Since the width of the meter's probe was approximately 3.5 cm (1.4 in.), the thermal meter recorded a centerline velocity. Total volumetric flow was calculated by multiplying the product of the measured gas velocity and the cross-sectional area of each stack by a correction factor to convert the centerline velocity in the stack to an estimated average velocity in the stack, accounting for the change in velocity profile from friction near the stack walls and accounting for the cross sectional area of the stack obstructed by the flow meter (see Appendices Section B.2). In some well configurations (31 of the 107 wells on which measurements were made), measurement through a temporary stack on the atmospheric tank was not technically feasible. In these cases, measurements were made by inserting a segment of pipe (with the thermal gas mass flow meter in the pipe) into the process line between the well and the atmospheric tank in order to measure the flow into the tank.

The methane fraction of the vented gas was assumed to be equal to the methane fraction in the normally produced gas. This was presumed to be a more accurate indicator of total methane emissions than measurements of the gas composition made through the temporary stack. The gas exiting through the temporary stack during the unloading period is a combination of the unloaded gas from the well and the gas initially in the vapor space of the tank (typically much lower in methane than the site's produced gas). At the end of the unloading, the tank will contain more methane, from the unloading, than was in the tank at the start of the unloading. This methane, which is associated with the

unloading event, will eventually be released as part of normal tank operations. Multiplying the measured vented gas volume by the methane fraction of the produced gas captures these emissions that occur because of the unloading but that are not released during the period when the tank is actively venting.

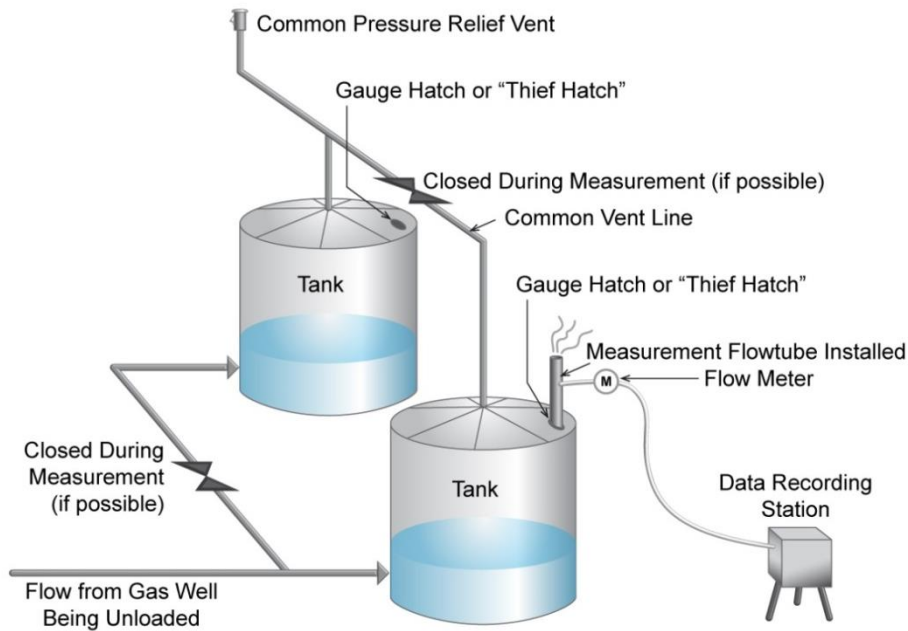


Figure 4-1. Conceptual diagram of a tank layout on a well site and the positioning of the temporary stack used to measure volume of gas vented during a liquid unloading

Uncertainty in these measurement methods is estimated at 10-20% of the measured emissions and this estimate is dominated by the assumed uncertainty in the flow, which includes both uncertainties in the stack gas volumetric flow measurement, and determining when flows return to zero. Variability in the gas composition from the well is expected to be much less than 10%. As described in the Results section, these measurement uncertainties are small compared to the combined sampling and measurement uncertainty, which are 50% or more of measured emissions.

Liquids can also be unloaded from a well using a plunger lift system. This liquid recovery operation holds a plunger at the top of the well, and either manually or by

automation occasionally closes (shuts-in) the well and releases the plunger, allowing it to fall down the well bore below the accumulated liquids. The well is then reopened, allowing the gas to push the plunger and the liquid back up the well bore as a slug of liquid. If the plunger returns to the top and the liquid and gas flow to the separator, there is no venting and all gas from the separator is routed to sales. In some cases, if the plunger does not return to the surface as expected, the plunger controller may bypass the separator and direct the flow to an atmospheric pressure vent, such as a vented tank. Directing flow to the lower pressure vent causes the plunger to return to the surface but also allows gas to vent. Plunger cycles may be initiated manually, on a timed interval, or based on certain well parameters such a reduced gas flow. In this work, measurements were made on both wells in which the unloading was automated through use of a controller (automatically triggered), and wells in which the plunger lift cycle was manually initiated by an operator (manually triggered).

In both the manually triggered plunger lift unloadings and the automatically triggered plunger lift unloadings, the volume of vented gas was measured using the same procedures as used for the wells without plunger lifts. For the automatically triggered unloadings, the measurement equipment was typically left in place for one to several days, making measurements continuously. This allowed automated plunger unloading events to be measured only when and if they occurred in routine operation, without artificially triggering the events. For all the plunger lift unloading events, the composition of the vented gas was assumed to be the same as the composition of the gas produced by that well. Produced gas composition was provided by site operators.

4.2.2 Sampling Strategies

Emission estimates reported through the EPA 2012 GHGRP [2] indicate that a small fraction of wells, in particular geographical regions, account for a large fraction of emissions from liquid unloadings. The sampling strategy employed in this work was to

sample most extensively in regions that were likely to dominate emissions (Appendices, Section B.1). Details of the sampling approach are provided in the Appendices. Briefly, the sampling team would visit a region for one or multiple weeks and sample a randomly selected subset of those wells that were unloading during that period. Consequently, more samples were collected on wells that unloaded more frequently. The features of these sample collection methods (preferential sampling in regions with high estimated emissions from unloadings and sampling of wells that tended to have high unloading frequencies) are important to consider when the data presented in this work are used to establish national emission estimates.

4.3 Results and Discussion

Methane emissions from liquid unloadings were measured at 107 natural gas wells. A summary of the geographical locations of the wells sampled is provided in Table 4-1.

For the 32 wells without plunger lifts (manually unloaded) sampled in this work, one event was typically sampled for each well; a few wells had more than one event sampled and for these wells, average values are reported. The unloadings of wells without plunger lifts sampled in this work had durations that lasted between 0.17 and 4.5 hr, and vented methane volumes that ranged between 550 and 135,000 standard cubic feet (scf) of methane per event (0.011 – 2.6 Mg). Representative time series for the methane emissions from wells without plunger lifts are shown in Figure 4-2. These three events shown had durations that ranged from 2.72 to 3.75 hr. Vented volumes for these three events shown ranged between 21,000 and 135,000 scf methane (0.40 – 2.6 Mg). As illustrated by these representative time series, some manual unloadings without plunger lift rapidly rose to a high flow rate, then maintained a steady flow throughout the event; others rose more slowly to a peak flow, then had variable flow during the event; still others rapidly rose to a peak flow, then had declining flows throughout the event. This

complex flow behavior makes it difficult to generalize about the flow characteristics for manually triggered unloadings of wells without plunger lifts.

Table 4-1. Unloading events measured in this work. Wells with manual unloadings typically had one event per well, while automated plunger lift unloadings had multiple events per well, a mapping of region boundaries is provided in the Appendices.

Type of Well	Initiation System	Wells with Unloadings sampled				
		US Total	Appalachian	Rocky Mountain	Gulf Coast	Mid-Continent
Plunger	Auto	25	0	20	1	4
	Manual	50	7	29	1	13
Non-Plunger	Manual	32	4	2	14	12
Total		107	10	51	16	29

For the 50 plunger lift wells with manually triggered unloadings, one event was typically sampled for each well. The manual unloadings of wells with plunger lifts sampled in this work had durations that lasted between 0.03 hr and more than 3 hr, and had vented methane volumes that ranged between approximately 200 and 49,000 scf methane per event (0.004 – 0.94 Mg). Representative time series for the methane emissions from a manual unloading are shown in Figure 4-3. These three events shown had durations that ranged from 1.2 to 40 min. Vented volumes ranged between 1,220 and 27,000 scf methane for the three events shown (0.02 - 0.52 Mg). As illustrated by these representative time series, some manually triggered unloadings with plunger lift rapidly rose to a high flow rate, then almost immediately fell rapidly, leading to a relatively short duration event; others rose rapidly to a peak flow that was maintained for 5-10 minutes or more; still others had complex flow patterns over an event lasting 10 minutes or more. As was the case for manually triggered unloadings without plunger lifts, this complex flow behavior makes it difficult to generalize about the flow characteristics.

For automated plunger lift wells, the sampling equipment was left in place for one to several days at each well, and typically more than one event was sampled for each well.

The automatically triggered unloadings (with plunger lift) sampled in this work had durations that lasted between <1 minute and more than 20 minutes, and vented methane volumes that ranged between 50 scf methane and more than 8,000 scf methane per event) (0.001 – 0.15 Mg). The numbers of events sampled per well ranged from 2 to more than 70; average values of emissions per event were used when multiple events were recorded. Representative time series for the methane emissions from automated plunger lift unloadings are shown in Figure 4-4. Individual unloading events for these three wells had durations that ranged from 2 to 20 min. Vented volumes per event ranged between 60 and 8600 scf methane for the three wells shown (0.001 – 0.15 Mg). As illustrated by these representative time series, some plunger lift wells with automated unloadings had emissions per event that were quite similar, and that occurred with a regular frequency. In contrast, however, some automated plunger lift wells had events that had qualitatively different emissions and/or variable event frequencies.

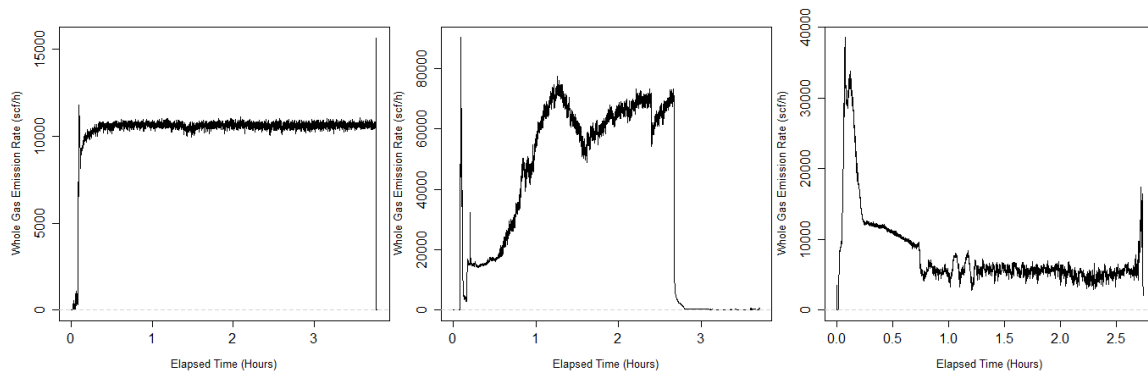


Figure 4-2. Representative time series of methane flow rates during manually triggered liquid unloadings from wells without plunger lifts (USH-47-0201 *left*; USH-47-0701 *middle*; UCG-03-0301 *right*); Note differences in horizontal and vertical scales.

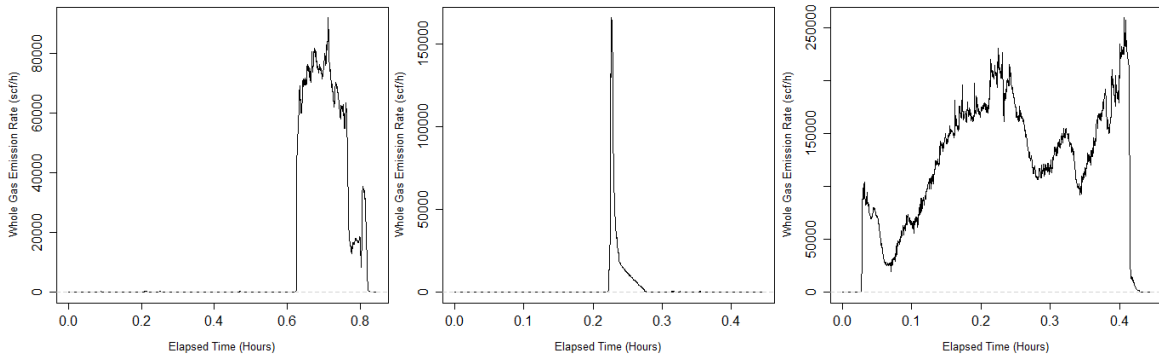


Figure 4-3. Representative time series of methane flow rates during manually triggered liquid unloadings from wells with plunger lifts (UBB-45-0101 *left*; UJR-46-0601 *middle*; USH-45-0202 *right*); Note differences in horizontal and vertical scales.

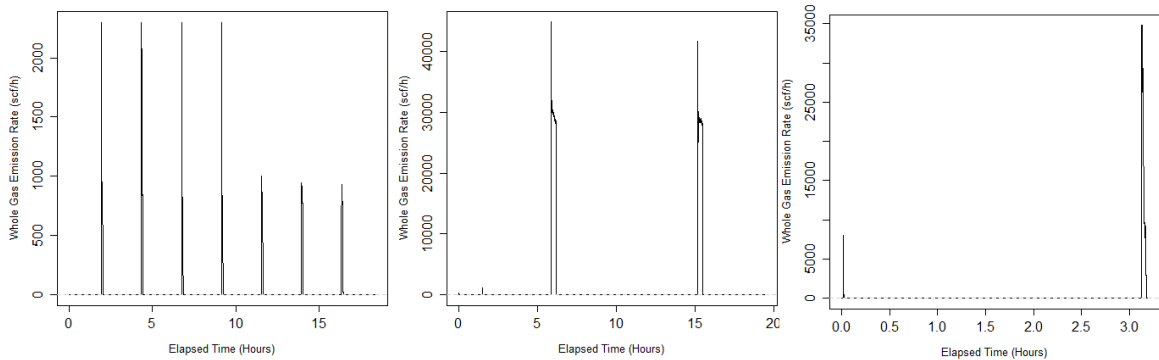


Figure 4-4. Representative time series of methane flow rates during automatically triggered liquid unloadings from wells with plunger lifts (UBB-42-0401 *top*; UBB-42-0201 *middle*; UEF-49-0501 *bottom*); Note differences in horizontal and vertical scales.

The Appendices (Section B.3) provides details of the unloading emissions and well characteristics for each of the 107 wells sampled in this work. A summary is provided in Figure 4-5. A relatively small number of wells have high emissions and most wells have much lower emissions. For example, 20% (6 of 32) of the wells account for 83% of the annual emissions for wells without plunger lifts, where annual emissions are estimated by multiplying the emission for an unloading event, measured in this work, by the number of times that well unloaded during calendar year 2012 or 2013 (whichever was the most recent report available), as reported by the well operator. The 6 wells that account for 83% of the annual emissions of wells without plunger lifts vent 6% of their collective

annual production. For manually and automatically triggered plunger lift wells, 20% of the wells account for 65% and 72% of the annual emissions, respectively. These wells vent 2% and 20% of their collective annual production, for manually and automatically triggered wells, respectively.

Because the sample distributions are not normally distributed about a mean, uncertainties in the average values are reported based on the results of a bootstrapping method, rather than as a simple standard deviation of the dataset. In the bootstrapping procedure, the original data set of each type of well was recreated by making random event selections, with replacement, from the data set. A total of 1000 of these re-sampled data sets were created and the mean value of the emissions for each re-sampled data set was determined. The 95% confidence intervals for the emission estimates represent the 2.5% and 97.5% percentiles of the means in the 1000 re-sampled datasets. So, for example, for the 25 automatically triggered plunger lift wells, a mean value for emissions per event, for each well, was calculated by selecting 25 emission measurements, at random and with replacement. This mean was tabulated and the process was repeated 1000 times to generate 1000 mean values. The 2.5% and 97.5% percentiles were determined to be 538 and 2,085 scf methane per event, and these values are the 95% confidence bounds for the mean value of the measurements (in this case 1,210 scf methane/event). The bootstrapping procedure leads to a combined sampling and measurement uncertainty. This uncertainty has a much larger range (typically 50% or more of the mean value, see Figure 4-5) than would be estimated from the uncertainty associated with the measurement alone (approximately 10-20% of the measured value) and is a reflection of the heterogeneity of well characteristics in the data sets and the underlying population of wells with unloading emissions.

Statistical analyses were conducted to identify well and unloading event characteristics that could explain the variability in the measured emission data. Variables that were considered included well pressures, well bore volumes, well ages, unloading event durations and unloading frequencies. The variable that explained the largest amount of

variability in the observed annual well emissions was unloading frequency, although there was also a positive correlation of event frequencies with well age (older wells had more unloading events per year than younger wells) and a negative correlation of annual emissions with well depth (deeper wells, which were generally newer, had lower annual emissions than shallower, generally older, wells). Correlations with emissions per event were generally weaker than for annual emissions. Additional details are provided in the Appendices (Section B.4). As shown in Figure 4-5, for wells without plunger lifts, average emissions for individual unloading events range between 21,000 and 35,000 scf methane per event, if the events are binned into wells that have less than 10 events per year, between 11 and 50 events per year, and between 51 and 200 events per year. The differences in annual emissions from manually unloaded wells without plunger lifts are largely due to the frequency of events, rather than the volume of gas emitted per event. For wells with plunger lifts, Figure 4-5 reports average emissions in two frequency bins. Manually triggered plunger lift wells were binned as a single group; all had less than 100 events per year (maximum observed value of 52 events per year). Automatically triggered plunger lift wells were also considered as a single category since all of these wells had more than 180 unloadings per year (average of 1,870 unloadings per year in the sampled population). Plunger lift wells that were manually triggered had average emissions per event of 9,450 scf methane. Plunger lift wells with automated triggering of the unloading had average emissions of 1210 scf methane per event.

The measured emissions per event were compared to predictions made using emission estimation methods commonly used in EPA GHGRP reporting. For wells with plunger lifts, for which the data required to make the estimates were available (66 wells), the emission estimates averaged 20,200 scf/event as compared to an average of 7,100 scf/event for measurements. The difference is statistically significant ($p = 0.0001$). In contrast, for wells without plunger lifts, the observations were higher (statistically significant difference) than the estimates. See Appendices (Section B.4) for more details.

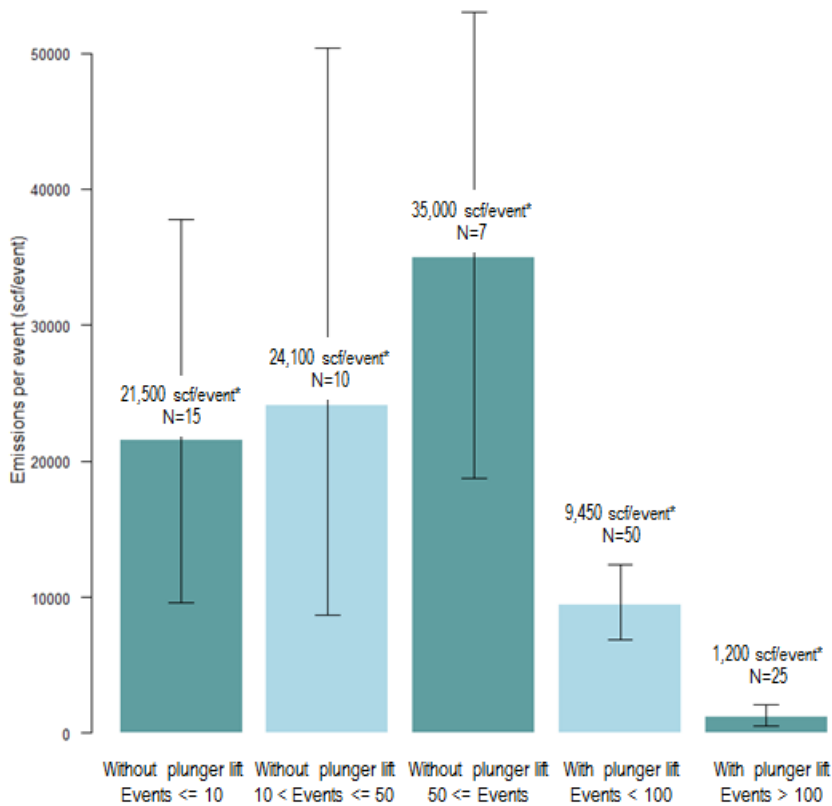


Figure 4-5. Average emissions per event for wells with and without plunger lifts, sorted by frequency of events (events per year per well)³.

4.4 Implications for National Emission Estimates

National emissions, based on the measurements made in this work, are estimated by multiplying an emission factor, based on the measurements, by an activity factor. Emission event counts, stratified into categories based on emission events per year per well, were chosen for the activity factor because of the process used for selecting wells to

³ *95% confidence bounds are:

	mean (scf methane/vent)	95% confidence bounds
Wells without plunger lifts ≤10 events per year:	21,500	9,600-37,800
Wells without plunger lifts 10<events≤50 per year:	24,100	8700-50,400
Wells without plunger lifts 50<events≤150 per year:	35,000	18,700-53,000
Wells with plunger lifts ≤100 events per year:	9,450	6,900-12,400
Wells with plunger lifts >100 events per year:	1,200	500-2,100

be sampled and because annual emission estimates for wells with unloadings depended most strongly on event frequency.

As documented in the Appendices (Section B.1), the measurement team typically visited production Basins for approximately a week, and sampled randomly selected wells that had scheduled (for manually triggered wells) or anticipated (for automatically triggered wells) unloading events for that week of sampling. This meant that the study team was far more likely to sample wells that unloaded weekly or more frequently, rather than wells that unloaded just a few times per year. This sampling approach resulted in a representative distribution of events, but not a representative distribution of wells. For example, 85% of the wells, without plunger lift, that have unloading emissions and that are operated by the companies that provided sampling sites in this work, had fewer than 10 emission events per year (See Appendices, Section B.5). In the measurements performed for this work, 15 of the 32 wells without plunger lift (47%) had 10 or less events per year. These wells are therefore under-represented in the measurement data, relative to their presence in the participating companies' overall well population. Because of differences in the distributions of event frequency between the sampled wells and the national population of wells, it would not be appropriate to choose an emission factor of emissions per well per year and an activity factor of number of wells, without adjusting for this difference in event frequency distribution.

An additional reason for stratifying wells by frequency of events in the activity factor is the data shown in Figure 4-5, which indicate a reasonable degree of consistency in per event emissions. Wells without plunger lifts had mean values of 21,000-35,000 scf methane/event. Wells with plunger lifts had mean values of 1,000-10,000 scf methane/event, but much larger ranges of event frequencies. For the calculations reported in this work, national, rather than regional averages of emissions per event will be used, due to the limited number of observations in individual regions.

In this work, national estimates of numbers of unloading events were based on a survey of the participating companies (see Appendices, Section B.5). Data on event counts from EPA GHGRP were not used since event counts for plunger and non-plunger wells are either partially reported or of uncertain quality. The national event counts were assumed to have the same proportion as reported by in the participant survey. Based on this survey, it was estimated that the 32,225 wells with plunger lifts (based on data from the 2012 GHGRP) have a total of 6.8 million events per year. Only 206,500 of these 6.8 million events are associated with wells that vent less than 100 times per year. Total annual emissions from plunger lift wells are estimated at 9.9 billion cubic feet of methane per year (bcf/yr) (190 Gg/yr), with 80% of those emissions associated with wells that vent more than 100 times per year (additional details in Appendices, Section B.5). For wells that vent more than 100 times per year, the average emissions per well per year are 1,400,000 scf per well per year (27 Mg/yr) with 95% confidence bounds of 600,000 – 2,500,000 scf (10 - 50 Mg, based on the confidence bounds in the emissions per event).

For wells without plunger lifts, it was estimated that 26,438 wells (based on data from the 2012 GHGRP) vent a total of 177,000 times per year, with total emissions of 4.4 bcf/yr (84 Gg/yr). Again, the wells that vent with highest frequency have the highest emissions per well. The 1.1% of wells that vent more than 50 times per year have average emissions of 3.2 million scf/yr. For wells without plunger lifts, however, these wells account for only 1.1% of the wells with unloading emissions, so the emissions from these wells venting at high frequency account for only 25% of emissions from wells without plunger lifts. Additional details are provided in Appendices, Section B.5).

The overall emission estimate for liquid unloadings (plunger and non-plunger wells) is 270 Gg (14 bcf/y), which is within a few percent of the national emissions estimated in either the 2012 GHG NEI (273 Gg/yr) or the 2012 GHGRP (276 Gg/yr). The 95% confidence range for this estimate is 190 – 400 Gg/yr, based on the reported confidence ranges in the per event emission factors, but not accounting for uncertainties in event counts. Appendices (Section B.5) reports sensitivity analyses that suggest uncertainties in

event count estimates may be large, which could have a significant impact on national emission estimates. Regardless of the exact national total of emissions, however, wells with high frequencies of unloadings have annual emissions that are a factor of 10 or more greater than the annual emissions of wells with low frequencies of unloadings.

4.5 Conclusions

Methane emissions from liquid unloadings were measured at 107 wells in natural gas production regions throughout the United States. Liquid unloadings clear wells of accumulated liquids to increase production, employing a variety of liquid lifting mechanisms. In this work, wells with and without plunger lifts were sampled. Most wells without plunger lifts unload less than 10 times per year with emissions averaging 21,000-35,000 scf methane (0.4-0.7 Mg) per event (95% confidence limits of 10,000-50,000 scf/event). For wells with plunger lifts, emissions averaged 1,000-10,000 scf methane (0.02-0.2 Mg) per event (95% confidence limits of 500-12,000 scf/event). Some wells with plunger lifts are automatically triggered and unload thousands of times per year and these wells account for the majority of the emissions from all wells with liquid unloadings. If the data collected in this work are assumed to be representative of national populations, the data suggest that the central estimate of emissions from unloadings (270 Gg/yr, 95% confidence range of 190-400 Gg/yr) are within a few percent of the emissions estimated in the EPA 2012 Greenhouse Gas National Emission Inventory (released in 2014), with emissions dominated by wells with high frequencies of unloadings.

4.6 References

(1) U.S. Environmental Protection Agency (EPA). Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2012, April 2014.

(2) U.S. Environmental Protection Agency (EPA). Greenhouse Gas Reporting Program, Data for Reporting Year 2012, 2013.

(3) American Petroleum Institute and America's Natural Gas Alliance (API/ANGA) Characterizing Pivotal Sources of Methane Emissions from Natural Gas Production,: Summary and Analyses of API and ANGA Survey Responses, Final Report, updated September, 2012, available at: <http://www.api.org/news-and-media/news/newsitems/2012/oct-2012/~~/media/Files/News/2012/12-October/API-ANGA-Survey-Report.pdf>

(4) Allen, D.T., Torres, V.M., Thomas, J., Sullivan, D., Harrison, M., Hendler, A., Herndon, S.C., Kolb, C.E., Fraser, M., Hill, A.D., Lamb, B.K., Miskimins, J., Sawyer, R.F., and Seinfeld, J.H. Measurements of Methane Emissions at Natural Gas Production Sites in the United States, *Proceedings of the National Academy of Sciences*, 110, 17768-17773, doi: 10.1073/pnas.1304880110 (2013).

5 ANALYSIS OF METHANE EMISSIONS ATTRIBUTABLE TO NATURAL GAS PRODUCTION

5.1 Introduction

A number of studies have examined greenhouse gas footprints of natural gas supply and use chains, and many of these studies have compared natural gas footprints to the greenhouse gas footprints for the production and use of other fuels [1-6]. In performing comparisons between natural gas and other fuel systems, it is important to recognize that parts of the natural gas supply chain, particularly production operations, produce both natural gas and liquid products. Co-production of natural gas and liquid products is particularly common in some of the most rapidly growing shale gas production regions, such as the Eagle Ford region in Texas [7]. When both gas and liquid products are generated by gas wells, the emissions of greenhouse gases from this part of the supply chain should be allocated to both gas and liquids production.

Recently Allen, *et al.*[8-9] have reported measurements of methane emissions from natural gas production sites, including wells that produced only gas and wells that produced both gas and liquids. All of these wells were classified as gas wells, all were in shale formations, and all were hydraulically fractured. Methane was the focus of these measurements because emissions of methane, a potent greenhouse gas, can significantly impact the supply chain greenhouse gas footprints of natural gas, relative to other fuels. Activities sampled included well completion flowbacks, liquid unloadings, pneumatic device operation, and equipment leaks from wells in routine operation. These data can be used to develop methane emission footprints of the production portion of the natural gas supply chain, with emissions allocated to the multiple products from the wells. For each of the wells sampled by Allen, *et al.* [8-9], gas composition, gas production and liquid production data are available [9]. This work reports allocations of methane emissions to natural gas and other gas and liquid products.

5.2 Methods

Allocation methods are commonly used in the analysis of emissions from supply chains when processes produce multiple products. For the case of a well that produces hydrocarbons that will eventually be separated into pipeline quality natural gas, natural gas liquids, and liquid hydrocarbon products, emissions from devices that handle all the products (e.g., a separator), should be allocated among multiple products. The most commonly used allocation methods are based on energy, mass and value [10-12].

In an energy based allocation, a well that produces 6000 standard cubic feet (scf) of gas for every barrel (bbl) of hydrocarbon liquid would generate equal amounts of energy as gas and liquid products, assuming a heating value of 1000 BTU/scf for gas and 6 million BTU/bbl for hydrocarbon liquid. For this simple example, if an emission allocation is based on energy, half of the emissions from the well would be assigned to the gas and half to the liquid product. In contrast, a mass based allocation, for the same simple example, would allocate 60% of the emissions to the gas and 40% to the oil, based on a gas density of 25 g/scf and an oil density of 100 kg/bbl. A value based allocation, based on prices of \$3.50 per thousand scf (Mscf) for gas and \$90 per bbl for hydrocarbon liquid, would assign 19% of the emissions to gas and 81% to the liquid.

The gas leaving a well site will typically contain quantities of ethane, propane, butane, and heavier hydrocarbons that are largely removed from the methane in the gas before the product is marketed as natural gas. In this work, emissions from well sites will initially be allocated to liquid products and to specific gas phase molecular species (methane, ethane, propane, butanes and pentane and heavier hydrocarbons). The emissions allocated to the gases will then be attributed to three main products: (1) *salable natural gas* (also known as dry natural gas, referring to the remaining gas once the liquefiable hydrocarbon portion has been removed)[13]; the composition of salable gas is based on commonly used life cycle analysis datasets for natural gas: 92.8% methane, and up to 5.54% non-methane hydrocarbon gases (the rest is N₂, CO₂, H₂S, and H₂O)[14] (% mass)), (2) *natural gas liquids*, which will be assumed to be the remainder of the

hydrocarbon gas leaving the well, and (3) *Hydrocarbon liquids (oil)* for which methane emissions will be assigned as a mixture.

Based on the assumed composition for *salable natural gas*, if the gas produced at each site has a non-methane hydrocarbon content that would produce a *salable natural gas* of up to 5.54% non-methane hydrocarbon (mass), all non-methane hydrocarbons are considered as part of salable natural gas (for that site there would be no natural gas liquids production). On the other hand, if the gas produced at each site is wetter (>5.54% non-methane hydrocarbon gases), non-methane hydrocarbon gases are assigned to *salable natural gas*, starting with ethane and then adding subsequent heavier hydrocarbons, until the 5.54% threshold is reached, the remaining portion of the gas leaving the well is considered *natural gas liquids*.

Properties and assumed economic values for each of these materials are provided in Appendices (Section C.1), Appendices also includes an expanded explanation of how non-methane hydrocarbon gases are split between salable natural gas and natural gas liquids (Appendices, Section C.4). Calculated across all sites, the weighted average composition (% mass) of salable natural gas is 97.3% methane, 2.61% ethane, 0.120% propane, 0.007 butanes, and 0.003% pentane and heavier hydrocarbons. For Natural Gas Liquids, the weighted average composition is 38.6% ethane, 27.8% propane, 18.8% butanes, and 14.8% pentane and heavier hydrocarbons (% mass).

For the case of the energy based allocation, a single heating value of 1,027 BTU/scf (higher heating value) was assigned to salable natural gas (Appendices, Section C.1), this heating value is commonly assumed in life cycle datasets [5, 14]. Nonetheless, a similar value is obtained when the mass weighted average heating value of salable natural gas across all sites is calculated (1,022 BTU/scf), based on the gas composition. For Natural Gas Liquids, an average heating value is derived for each site, based on its particular Natural Gas Liquids composition. (Mass weighted average heating value of natural gas

liquids across all sites is 2,349 BTU/scf (higher heating value)) (See Appendices, Section C.5).

Emissions for each product were allocated based on mass, energy and value, for each product (salable natural gas, natural gas liquids, and oil), for each of the individual sampling sites that were reported by Allen, *et al.* [8-9]. Average normalized emissions for each product are reported as the sum of emissions over all sites attributed to a particular product, divided by the sum of production of that product, over all sites. So, for example, the emissions attributed to natural gas liquids would be summed over all sites then divided by the production of natural gas liquids, summed over all sites.

Production for the sites can be reported either as instantaneous production rate at the time of the measurement or as ultimate production. In this work, the production rate will be reported as an estimated ultimate recovery, over 30 years, for each well. The ultimate recovery is estimated for each well using the age of the well at the time of the measurement, the instantaneous production at the time of the measurement, and an assumed production decay rate over 30 years of production, as described in the Appendices (Section C.2). Sensitivity analyses, using instantaneous production rates and shorter well life, are also reported.

Results in this work are reported as a ratio. For emissions from wells in routine production, the numerator in the ratio is total estimated emissions, over 30 years, based on a measured instantaneous emission rate and an assumption that emissions (except for well completions) remain constant over 30 years. The denominator is the estimated ultimate recovery from the well, over 30 years. For completion events, which are assumed to only occur once over the life of a well, the numerator is the emissions from the single event. The denominator is the estimated ultimate recovery from the well, over 30 years.

5.3 Results and Discussion

5.3.1 Emissions from wells in routine production

Table 5-1 reports emissions from pneumatic controllers, pneumatic pumps, and leaks from equipment sampled by Allen, *et al.* [8-9]. Allocated emissions for individual sites are shown in Appendices, Sections C.3-C.6. To arrive at total emissions for each site, emissions from leaks, pneumatic controllers and pneumatic pumps are summed. All leaks at sites were sampled, but not all controllers or pumps. For controllers and pumps, the average emission rate for each site was multiplied by the total number of devices reported for that site. Averages over multiple sites are taken on a production weighted basis, as described in the Methods section. Emissions assigned to each product are summed over all sites and divided by the production rate of that product, summed over all sites.

Table 5-1. Ratio of methane emissions allocated to each product (Natural Gas, Natural Gas Liquids, and Oil) divided by its respective production, based on a **mass** allocation, **energy** allocation, and **price** allocation. The values represent the ratio of the sum of emissions divided by the sum of production across all sites where each specific product is produced. From a total of 150 production sites, 144 sites reported natural gas production (for 3 sites production data was reported as “not available”; three sites reported zero gas production), 39 sites reported oil production, and 51 reported Natural Gas Liquids production.

	Pneumatic Controllers	Pneumatic Pumps	Equipment Leaks	Total
MASS ALLOCATION				
Emitted Methane (scf) / Produced Natural Gas (scf)*	0.0008	0.0002	0.0002	0.0011
Emitted Methane (Mg) / Produced Natural Gas Liquids (Mg)**	0.0014	2.2E-05	0.0001	0.0016
Emitted Methane (Mg) / Produced Oil (Mg)	0.0011	1.9E-06	0.0001	0.0012
ENERGY ALLOCATION				
Emitted Methane (scf) / Produced Natural Gas (scf)*	0.0008	0.0002	0.0002	0.0012
Emitted Methane (Mg) / Produced Natural Gas Liquids (Mg)**	0.0013	2.0E-05	0.0001	0.0015
Emitted Methane (Mg) / Produced Oil (Mg)	0.0010	1.7E-06	0.0001	0.0012
PRICE ALLOCATION				
Emitted Methane (scf) / Produced Natural Gas (scf)*	0.0007	0.0002	0.0002	0.0011
Emitted Methane (Mg) / Produced Natural Gas Liquids (Mg)**	0.0017	4.6E-05	0.0002	0.0019
Emitted Methane (Mg) / Produced Oil (Mg)	0.0016	3.9E-06	0.0002	0.0017

**Produced Natural Gas* refers to *salable natural gas* (also known as dry natural gas, referring to the remaining gas once the liquefiable hydrocarbon portion, propane and heavier, has been removed [13], with the remaining non-methane hydrocarbon gases [14], being allocated to Natural Gas Liquids.

** *Produced Natural Gas Liquids, NGL* refers to the remaining non-methane hydrocarbon gases that are not part of salable natural gas. (For expanded explanation of how non-methane hydrocarbon gases are split between salable natural gas and natural gas liquids see *Appendices, Section C.4*).

If the instantaneous emission rate is normalized by instantaneous production rate at the time of the measurement, emissions from pneumatics and leaks from wells in routine production are 0.0003 scf of methane per scf of produced natural gas, compared to the 0.0011 – 0.0012 scf of methane per scf of produced natural gas reported in Table 5-1, when emissions over 30 years and an ultimate production based on a well lifetime of 30 years is assumed (see Appendices, Section C.8). The difference is due to the relatively young age of the wells in the data set (average age of wells, weighted by gas production is 1.3 years), and the assumption that emissions stay constant over the life of the well. If the assumed well life is reduced to 15 years, emissions are reduced by half but production is only reduced by, on average, 22% (see Appendices, Section C.7). This leads to an estimated natural gas emission rate of 0.0007 scf of methane per scf of produced natural gas.

5.3.2 Emissions from Completion Flowback Events

Allen, *et al.* [8-9] also made direct measurements from 27 well completion events. Production data are reported for 19 of these 27 wells; 12 report both gas and oil production, and 7 report production of gas only. Table 5-2 shows the average ratio of methane emissions from the 19 completions, divided by the total estimated ultimate recovery of each component, over 30 years, from the 19 wells. This average represents the sum of emissions divided by the sum of production across the 19 measured events. (Ratios and allocated emissions for each separate event are shown in the Appendices, Section C.9). Table 5-3 shows the sum of the average completion, pneumatic device and leak measurements. The contribution of completion events is small compared to the emissions from wells in routine production.

Table 5-2. Ratio of methane emissions from completion flowbacks allocated to specific gas phase molecular species (methane, ethane, propane, butanes and pentane and heavier hydrocarbons) and to each product (Natural Gas, Natural Gas Liquids, and Oil) divided by its respective production. Results are shown in mass, energy, and price basis.

	Mass Basis	Energy Basis	Price Basis
Emitted Methane (scf) / Produced Natural Gas (scf)*	1.8E-05	1.8E-05	1.1E-05
Emitted Methane (Mg) / Produced Natural Gas Liquids (Mg)**	3.5E-05	3.4E-05	4.2E-05
Emitted Methane (Mg) / Produced Oil (Mg)	1.8E-05	1.7E-05	2.6E-05

Table 5-3. Ratio of TOTAL methane emissions from completion flowbacks, pneumatic devices and leaks and routine emissions) allocated to each component (Natural Gas, Natural Gas Liquids, and Oil) divided by its respective production. Results are shown on mass, energy, and price bases.

	Mass Basis	Energy Basis	Price Basis
Emitted Methane (scf) / Produced Natural Gas (scf)*	0.0012	0.0012	0.0011
Emitted Methane (Mg) / Produced Natural Gas Liquids (Mg)**	0.0016	0.0015	0.0020
Emitted Methane (Mg) / Produced Oil (Mg)	0.0012	0.0012	0.0018

Emissions from pneumatic controllers, pneumatic pumps, leaks from equipment and well completion flowback events (categories that were directly measured by Allen, *et al* [8-9]., represent 0.11% - 0.12% (vol.) of the total natural gas production when emissions are allocated among natural gas, natural gas liquids and oil; For those same categories, the current EPA national greenhouse gas inventory (data for calendar year 2012, released in 2014) reports a total of 792 Gg of methane/yr or 0.14%, on a volume basis of total natural gas production (29.5 trillion cubic feet of natural gas produced in the US in 2012 [16].) when allocation to co-products is not considered.

Table 5-4 lists additional sources of methane emissions from natural gas production that are quantified in the EPA national greenhouse gas inventory, but were not measured or were measured in very small sample sizes by Allen, *et al.* [8-9]. Table 5-4 disaggregates these emissions, based on whether the equipment handles natural gas (NG) and natural gas liquids (NGL) alone (NG+NGL), oil alone, or all products from the well (NG + NGL + OIL)⁴. The rationale for the disaggregation of each source into the three products is shown in Section C.10 of the Appendices. For categories that allocate emissions to all three final products (NG + NGL + oil), 85.1% of the emissions from each category will be allocated to Natural Gas. This 85.1% is based on the ratio of the average mass content in natural gas to the mass in all well products, over all wells on which measurements were made. For source categories that allocate emissions to Natural Gas and Natural Gas Liquids only (NG + NGL), 94.1% of the emissions are allocated to natural gas. This value is based on the ratio of the average mass content in natural gas to the combined mass NG and NGL products, over all wells on which measurements were made. Combined, these mass based allocations for natural gas production sources not directly estimated by Allen, *et al.* [8-9] allocate an estimated 1055 Gg of methane emissions to natural gas product, which is 0.19% (volume) of the 28.5 trillion cubic feet of natural gas produced in the US in 2011. Since these national estimates capture a population of wells of all ages, it is assumed that the percentage of natural gas emitted (0.19%) would be the same for current emissions from all wells of all ages divided by current production of all wells of all ages or well lifetime emissions divided by EUR.

If these additional emissions are added to the allocated emissions from pneumatics, equipment leaks, and completion flowbacks described in this work (0.12% of natural gas production), the total methane emissions would represent 0.31% (vol.) of total natural gas

⁴For categories that allocate emissions to all three final products (NG + NGL + oil), 85.1% of the emissions from each category are allocated to Natural Gas (percentage based on the previously described energy basis allocation).

For categories that allocate emissions to Natural Gas and Natural Gas Liquids only (NG + NGL), 94.1% (percentage based on the previously described energy basis allocation) of the emissions from each category are allocated to natural gas.

production, a value that is roughly 26% lower than the 0.42% reported by Allen, *et al.* [8], not accounting for co-product allocation.

Table 5-4. Description of emission sources not directly measured by Allen et al. [8] that are considered in the EPA GHG national inventory, showing how the emissions are allocated to the products considered in this work. (NG = Natural Gas, NGL = Natural Gas Liquids). The column showing net emissions refers to total emissions from each category before the allocation to the corresponding products. Last row shows total emissions after allocation. (Rationale for the disaggregation of each category is provided in the Appendices, Section C.10.)

EPA GHG Inventory Activity	Net Emissions (Gg methane/yr)	Allocate emissions to		
		NG + NGL + Oil	NG + NGL	Oil only
Refractures	143	✓		
Gas wells without HF	13	✓		
Gas wells with HF	15	✓		
Separators	57	✓		
Meters/Piping	54	✓		
Heaters	18	✓		
Dehydrators	16		✓	
Workovers without HF	0.3	✓		
Liquids Unloading (without plunger lifts)	149	✓		
Liquids Unloading (with plunger lifts)	108	✓		
Kimray Pumps	185		✓	
Condensate Tanks without Controls	94			✓
Condensate Tanks with Controls	52			✓
Gas Engines	227		✓	
Dehydrators Vents	41		✓	
Small Reciprocating Compressors			✓	
Large Reciprocating Compressors	49		✓	
Large Reciprocating Stations			✓	
Pipeline Leaks	90		✓	
Well Drilling	0.4	✓		
Vessel Blowdowns	0.4		✓	
Pipeline Blowdowns	2		✓	
Compressor Blowdowns	2		✓	
Compressor Starts	3		✓	
Pressure Relief Valves	0.4		✓	
Mishaps	1	✓		
Total Emissions allocated to NG		475 Gg	580 Gg	

5.3.3 *Regional analysis*

Allen, *et al.* [8-9] reported measured emissions from natural gas production sites from four different regions in the United States: Appalachian (45 sites), Gulf Coast (58 sites), Mid Continent (23 sites), and Rocky Mountains (18 sites). Methane emissions (sum of emissions from pneumatics and equipment leaks) are also allocated on a regional basis (mass based allocation), with the intention of highlighting regional differences in emission rates as well as in total production, and the effects that the proposed co-allocation scheme would have on each region's emissions. Table 5-5 shows allocated methane emissions for each region.

Table 5-5. Methane emissions allocated to each salable product (Natural Gas, Natural Gas Liquids, and Oil), total estimated ultimate recovery for each product and ratio of emissions to production, based on a **mass** allocation. The values represent the ratio of the sum of emissions divided by the sum of production across all sites where each specific product is produced, considering a 30 year well lifetime and for each region. For each region, the number of sampled wells (within the sampled sites) is shown.

	Appalachian (164 wells)	Gulf Coast (146 wells)	Mid Continent (76 wells)	Rocky Mountains (78 wells)
Total Methane Emissions (Mg)	15,801	28,143	15,975	4,272
Methane emissions allocated to Natural Gas (Mg)	15,736	21,288	13,994	2,175
Methane emissions allocated to Natural Gas Liquids (Mg)	64	1,928	1,201	1,394
Methane emissions allocated to Oil (Mg)	2	4,927	780	703
EUR Natural Gas (Mg)	23,770,765	20,371,250	2,729,781	363,904
EUR Natural Gas Liquids (Mg)	38,492	2,505,534	138,572	265,674
EUR Oil (Mg)	1,037	5,014,564	114,540	210,754
Emitted Methane (scf) / Produced Natural Gas (scf)	0.0007	0.0010	0.0051	0.0060
Emitted Methane (Mg) / Produced Natural Gas Liquids (Mg)	0.0017	0.0008	0.0087	0.0052
Emitted Methane (Mg) / Produced Oil (Mg)	0.0019	0.0010	0.0068	0.0033

The Appalachians region shows the biggest difference between gas and oil production; where combined production for Natural Gas and Natural Gas Liquids is four orders of magnitude larger than the Oil production (for the measured wells). In the case of the Gulf Coast region and the Mid Continent region, combined NG and NGL production is just one order of magnitude larger than oil production, while for the Rocky Mountains, gas and oil production are of the same order of magnitude.

When the (measured) wells of all regions are considered (as analyzed in the previous section) (national average) combined production of Natural Gas (47,235,700 Mg) and Natural Gas Liquids (2,948,272 Mg) is one order of magnitude larger than the oil production (5,340,895 Mg); with a higher production of oil relative to Natural Gas Liquids. Consequently, a majority of the emissions (85%) are allocated to Natural Gas while 10% are allocated to Oil, and 5% to Natural Gas Liquids.

For the Appalachian region over 99% of the emissions are allocated to Natural Gas, with almost all the remaining emissions being allocated to natural gas liquids. In the case of the Gulf Coast region, 70% of the emissions are allocated to Natural Gas; from the remaining emissions, 18 percent is allocated to oil and 12% to Natural Gas Liquids. For the Mid Continent, 92% of the emissions are allocated to Natural Gas, with remaining emissions being allocated in similar amounts to Natural Gas Liquids and oil. The Rocky Mountains represent the region where the smallest fraction of emissions is allocated to Natural Gas (43%), with 32% of the emissions allocated to Natural Gas Liquids, and 25% to oil. This is driven by the similar mass content of the three products.

The ratios of methane emissions allocated to natural gas production divided by its production are 5 to 6 times higher for the Mid Continent (0.0051) and Rocky Mountains (0.0060) than for the other two regions (Appalachian: 0.0007, Gulf Coast: 0.0010).

Wells in the Appalachian and Gulf Coast regions have a similar EUR for natural gas; 148,000 Mg per well and 141,500 Mg per well, respectively. These values are considerably higher than for the Mid Continent, where per well For the Rocky Mountain region EUR for natural gas is 5,000 Mg per well. The higher ratios observed in the Mid Continent and Rocky Mountains are driven by a similar order of magnitudes of emissions across all regions but a significantly lower production per well

5.3.4 Comparison to other databases:

The National Energy Technology Laboratory (NETL) prepared a life cycle assessment (LCA) for natural gas [5], both for conventional and unconventional sources. Under the life cycle stage called “Raw Material Acquisition”, the source categories include emissions from well construction, well completions, liquid unloading, workovers, other point source emissions (gas released from wellhead and gathering equipment), other fugitive emissions, and valve fugitive emissions. Data for the LCA comes primarily from EPA emission factors and inventories, which have been subject to uncertainty due to the lack of direct measurements. Estimates are based on the EPA GHG inventory released in 2011, which were higher than the most recent EPA estimates, released in 2014 [16].

NETL reports 0.366 kg of methane emitted per Million BTU of natural gas extracted or 0.0196 scf methane/ scf of extracted natural gas (NETL uses a natural gas energy content of 1,027 BTU per cubic feet of natural gas, the same value used in the present work (Appendices, Section C.1)), from the raw material acquisition stage from shale gas wells. This emission rate is an order of magnitude higher than the value presented in this work (0.0031 scf of methane emissions over the life of a well allocated to natural gas/scf of produced natural gas over a well lifetime) and higher than the value reported by Allen, et al. (0.0042 scf current emissions of methane/scf of current natural gas production) for unallocated emissions.

Argonne National Laboratory prepared a LCA comparing shale gas to conventional natural gas [15]. The report is based on EPA estimates and emission factors, and acknowledges the uncertainty in EPA’s estimates and modifications and revisions. Estimates are based on the EPA GHG inventory released in 2011 which were higher than the most recent EPA estimates, released in 2013 [16]. Based on EPA’s revisions to the inventory, as well as recently released reports of measurements of methane emissions, such as Allen, *et al.*[8], Argonne updated their inventories [17].

Argonne's GREET (Greenhouse gases, Regulated Emissions, and Energy use in Transportation) model is used for the analysis, developing a specific pathway for shale gas. The production stage includes emissions from well completions, workovers, liquid unloading, and well equipment (field separation equipment, gathering compressors, normal operations, condensate collection, compressors venting and upsets). The current inventory estimates a total of 120.7 g of methane per million BTU of natural gas, or a total of 0.0064 scf of methane/scf of natural gas produced, for the production stage, which represents roughly 50% of the emissions estimated with the previous version of the GREET model, and which is a factor of three higher than the value reported in this work.

5.4 Conclusions

Mass, energy, and price basis allocation methods have been used to assign methane emissions from natural gas production to the three main products of production activity: salable natural gas, oil, and natural gas liquids. On a national scale, approximately 85% of the emissions from the well site are assigned to natural gas, but regional variability is observed. Methane emissions allocated to salable natural gas reported in this work are lower than those reported in commonly used LCA data sets.

5.5 References

- (1) Alvarez RA, Pacala SW, Winebrake JJ, Chameides WL, Hamburg, SP (2012). Greater Focus Needed on Methane Leakage from Natural Gas Infrastructure Proceedings of the National Academy of Sciences 109: 6435–6440.
- (2) Jaramillo P, Griffin WM, Matthews HS (2007). Comparative lifecycle air emissions of coal, domestic natural gas, LNG, and SNG for electricity generation. Environ. Sci. Technol. 41: 6290–6296.

- (3) Laurenzi IJ, Jersey GR (2013). Life Cycle Greenhouse Gas Emissions and Freshwater Consumption of Marcellus Shale Gas Environ. Sci. Tech. 47: 4896-4903.
- (4) Pacsi AP, Alhajeri NS, Zavala-Araiza D, Webster MD, Allen DT (2013). Regional air quality impacts of increased natural gas production and use in Texas. Environ. Sci. Tech. 47: 3521-3527.
- (5) NETL. (2011). Life Cycle Greenhouse Gas Inventory of Natural Gas Extraction, Delivery and Electricity Production. U.S. Department of Energy.
- (6) Venkatesh A, Jaramillo P, Griffin, WM, Matthews HS (2012). Implications of changing natural gas prices in the United States electricity sector for SO₂, NO_x, and life cycle of GHG emissions Environ. Res. Lett. 7: 034018.
- (7) EIA, (2013). What is shale gas and why is it important? Retrieved 9/25/2013 from http://www.eia.gov/energy_in_brief/article/about_shale_gas.cfm
- (8) Allen DT, Torres VM, Thomas J, Sullivan D, Harrison M, *et al.* (2013). Measurements of Methane Emissions at Natural Gas Production Sites, Proceedings of the National Academy of Sciences in press.
- (9) Allen DT, Torres VM, Thomas J, Sullivan D, Harrison M, *et al.* (2013). Measurements of Methane Emissions at Natural Gas Production Sites: Study Appendices and Database, Austin, Texas: University of Texas <http://dept.ceer.utexas.edu/methane/study/>
- (10) Babusiaux D, Pierru A (2007). Modelling and allocation of CO₂ emissions in multiproduct industry: The case of oil refining Applied Energy, 84: 828-841.
- (11) Ekvall T, Tillman A-M (1997). Open-Loop Recycling: Criteria for Allocation Procedures. The International Journal of Life Cycle Assessment, 2 (3): 155-162.

- (12) Wang, M., Hanjie, L., & Molburg, J. (2004). Allocation of Energy Use in Petroleum Refineries to Petroleum Products. *The International Journal of Life Cycle Assessment* , 9 (1), 34-44.
- (13) EIA. 2011. Glossary. Retrieved 8 12, 2013, from <http://www.eia.gov/naturalgas/annual/pdf/glossary.pdf>
- (14) NETL. 2012. Role of Alternative Energy Sources: Natural Gas Technology Assessment. U.S. Department of Energy.
- (15) Argonne National Laboratory. (2011). Life-Cycle Analysis of Shale Gas and Natural Gas . Oak Ridge, TN: U.S. Department of Energy.
- (16) EPA. (2014, April 15). Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2012. Retrieved 6 28, 2014, from <http://www.epa.gov/climatechange/ghgemissions/usinventoryreport.html>
- (17) Argonne National Laboratory. (2013). Updated Fugitive Greenhouse Gas Emissions for Natural Gas Pathways in GREET Model. Oak Ridge, TN: U.S. Department of Energy.

6 CRITERIA AND AIR TOXIC EMISSIONS FROM SHALE GAS PRODUCTION

6.1 Introduction

The analysis for criteria pollutants and air toxic emissions from shale gas production focuses on the Barnett Shale region in Texas, which has shown a significant growth in gas production activity during the last ten years, with production of 0.11 billion cubic feet per day in 2000, increasing to 5 billion cubic feet per day by 2011 [1,2]. Total natural gas withdrawals in the US in mid-2011 were approximately 70 billion scf/d (bcf/d); with approximately 22 bcf/d in Texas [3]. At 5 bcf/d of production, the Barnett Shale is one of the largest natural gas production regions in the US. The region includes 24 counties to the north and west of Fort Worth, with a total of more than 14,000 producing wells [4].

Since 2010, the Texas Commission on Environmental Quality (TCEQ) has deployed automated gas chromatographs that have recorded hourly averaged atmospheric concentrations of hydrocarbons in the Barnett Shale production region and neighboring areas. The TCEQ has also developed estimates of emissions from over 10,000 production sites in the Barnett Shale region. Figure 6-1 illustrates the locations of the monitoring sites that will be examined in this work, along with the inventoried VOC emissions from the production sites.

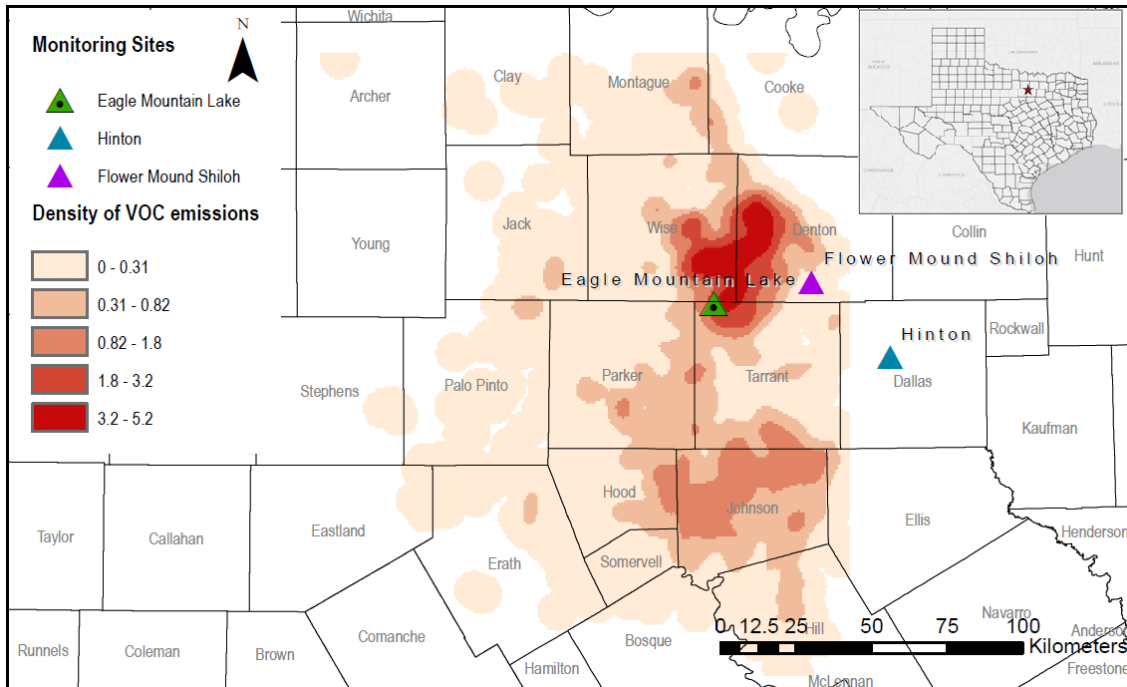


Figure 6-1. Density of total VOC emissions (tons per year per square kilometer) based on the locations of natural gas production sites. The map also shows the air monitoring sites (Eagle Mountain Lake (EML), Flower Mound Shiloh and Hinton). The density of emissions is calculated from roughly 8,000 sites which report 19,050 tpy of VOC emissions for the TCEQ Barnett Shale Area Special Inventory.

The goals of the analyses to be presented in this chapter are (1) to report diurnal, monthly and seasonal trends in VOC composition and magnitude; (2) to compare the magnitudes of observed concentrations to concentrations predicted based on the emission inventory using a Lagrangian air quality model; (3) to evaluate the effect that episodic emissions have on the overall predicted concentrations and (4) to assess the overall contributions of the shale gas emissions to benzene concentrations in the region.

6.2 Methods

6.2.1 *Ambient measurements of VOC*

Hourly concentrations for 46 non-methane volatile organic compounds at three sites will be examined in this work. These three sites were selected since they represent a site in the middle of the Barnett Shale (Eagle Mountain Lake, EML), a site at the periphery of the Barnett Shale (Flower Mound-Shiloh) and an urban site, relatively far from the Barnett Shale (Hinton). Eagle Mountain Lake (EML (32.9879°, -97.4772°)), is located in Tarrant County northwest of the Fort Worth urban area. As shown in Figure 6-1, it is located in approximately the geographical center of the Barnett Shale natural gas production region, so EML will serve as the core site for characterizing ambient VOC composition in the shale gas production region. The Flower Mound Shiloh site (33.0459°, -97.1300°) is located northwest of the Dallas urban area and north east of the EML site. Under typical wind conditions, with flow from the south, it is not as strongly influenced by natural gas production activities as the EML site. The Hinton site (32.8201°, -96.8601°), located in the Dallas downtown area, is used to contrast urban sites in the region to the sites influenced by natural gas production emissions. More details about these sites are available from the TCEQ [5].

Hydrocarbon measurements were made using automated gas chromatographs (auto-GCs), which measure concentrations of 46 hydrocarbons on an hourly basis using flame ionization detectors. The auto-GC instrument utilizes two capillary columns that separate light hydrocarbons and heavy hydrocarbons. Samples are collected for a 40 minute period by adsorbing hydrocarbons onto an automated thermal desorber; after the collection period, a 20 minute gas chromatography analysis is performed. Species concentrations determined by the auto-GC are reported as hourly averages based on average concentrations during the 40 minute sampling time. Concentrations are based on FID responses for propane and benzene. Propane is used to calibrate the column for lighter hydrocarbons, while benzene is used to calibrate the column for heavier hydrocarbons²⁴. Hydrocarbon concentration and wind data from the period of April 19, 2010 through

December 31, 2011 were examined; the starting date for the analyses is when the auto-GC measurements were initiated at EML.

6.2.2 VOC Emission inventory

The TCEQ reports VOC emissions from individual production sites in the Barnett Shale special inventory [6]. The Barnett Shale Area Special Inventory contains information on a total of 8,025 sites which report a total of 19,050 tons per year (tpy) of VOC emissions. Emissions are disaggregated into 18,466 point sources classified by type of source (92% of the VOC emissions report source type, the other 8% is site specific only). All sites have latitude and longitude reported. The TCEQ provided an emission calculator to assist companies with the emissions estimates [7]. The TCEQ requested emissions estimated with site-specific data. On-site measurements were preferred to emissions-factor based methods but were not required [4].

Table 6-1 shows the total VOC emissions by source type from the TCEQ inventory. Condensate tanks are the largest source of VOC emissions followed by fugitives, water tanks, and engines. VOC emissions from sites that did not report disaggregated sources are listed as without source type (8.1% of the total reported emissions).

Table 6-1. Total VOC emissions by source type in the Barnett Shale Special Inventory

Type of source	Total VOC emissions (tpy)	% of total VOC emissions
Amine unit	2.47	0.013%
Boiler	0.30	0.002%
Condensate tank	10,464	54.9%
Engine	749	3.93%
Flare	19.5	0.103%
Frac Tank	3.46	0.018%
Fugitives	3,935	20.7%
Glycol Dehydrator	65.8	0.345%
Heater	1.68	0.009%
Loading	511	2.68%
Oil Tank	221	1.16%
Other	9.47	0.050%
Separator	161	0.846%
Vent	84.3	0.442%
Water tank	1,286	6.75%
Without source type	1,535	8.06
Total	19,050	100%

As shown in Table 6-1, the TCEQ inventory predicts that VOC emissions will be dominated by tanks and that fugitives account for about 21% of VOC emissions. A recent field measurement campaign¹⁷ has found, however, that current emission factors may underestimate equipment leaks and emissions from pneumatic devices in the Barnett Shale. So, for a series of sensitivity analyses, modified emission factors for equipment leaks and emissions from pneumatic devices (which are both considered fugitives in the TCEQ inventory) will be used.

In addition, the inventory does not contain estimates of emissions from events such as well completions and liquids unloading. The potential impacts of these intermittent, short duration, and potentially large emission sources (episodic emissions) will be examined in a series of sensitivity studies, based source data recently collected on these types of events. [8]

6.2.3 Lagrangian air quality modeling

The AERMOD dispersion model was used to predict VOC concentrations at EML. AERMOD is a steady-state plume dispersion model that can be used to predict concentration distributions under a wide variety of conditions and in moderate to complex topography [9]. It uses a Gaussian distribution in the horizontal direction in the stable boundary layer. The AERMOD modeling system consists of one main program (AERMOD) and three pre-processors (AERSURFACE, AERMAP, and AERMET).

For input to the AERSURFACE pre-processor, National Land Cover data was obtained from the USGS. For AERMOD's meteorological preprocessor (AERMET), near surface characteristics as well as upper-air conditions are needed; these data were obtained from National Weather Service data and NOAA. Additionally, AERMET requires onsite meteorological data, which in this case was obtained from the measurements from the EML monitoring site. With these three datasets AERMET estimates planetary boundary layer parameters²⁹. The locations of point sources with known latitude and longitude were extracted from the Barnett Shale Area Special Inventory and were used as input for AERMAP, as well as the USGS 30 m National Elevation Dataset covering the studied region.

Four discrete AERMOD modeling domains were used to evaluate scenarios incorporating sources within 100, 50, 25, and 10 km radii from the EML monitoring site. Emissions from point sources within these distances, obtained from the Barnett Shale Area Special Inventory, their location and reported total VOC emissions (total VOC emissions with

known latitude and longitude), were used as inputs to the dispersion model, along with hourly meteorological data for the 20 month sampling period.

The model was run with just one receptor, corresponding to the location of the EML monitoring site; the various plumes generated at each of the thousands of sources are combined at this receptor point. Hourly total VOC concentrations were predicted for the EML site for the 20 month period.

6.3 Results and Discussion

6.3.1 Hydrocarbon concentration measurements

Figures 6-2a and 6-3a show measured diurnal and monthly average concentrations of hydrocarbons at the EML site. Light alkanes dominate, with ethane, propane and n-butane accounting for approximately 70% of the identified hydrocarbon concentrations, expressed as ppbC. This composition is consistent with the expected composition of emissions from natural gas production activities.

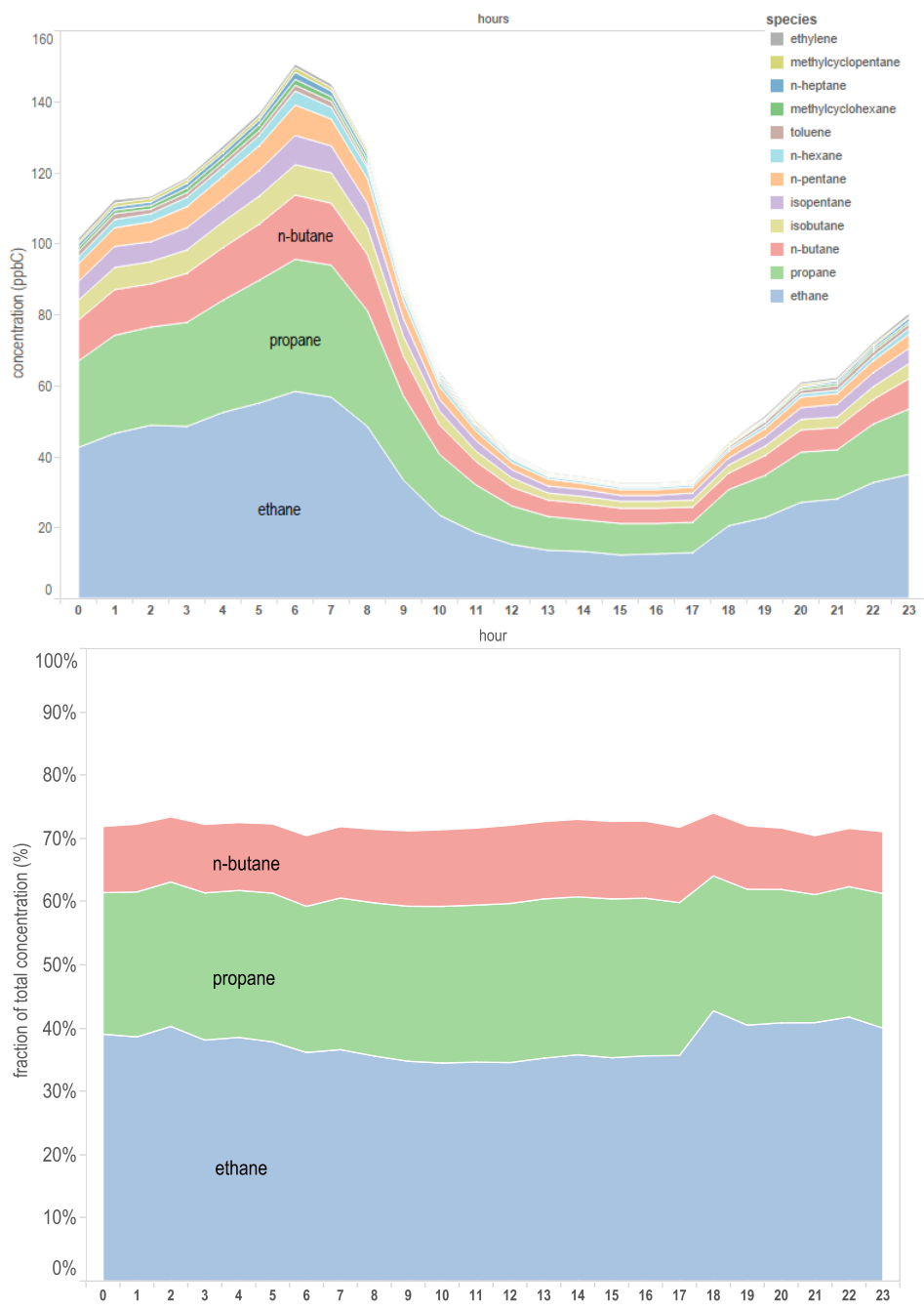


Figure 6-2. . (a) Average diurnal pattern of hydrocarbon concentrations (ppbC) at EML; data for each hour are averaged over 20 months of sampling. (b) Percentage of ppbC accounted for by ethane, propane and n-butane, at EML.

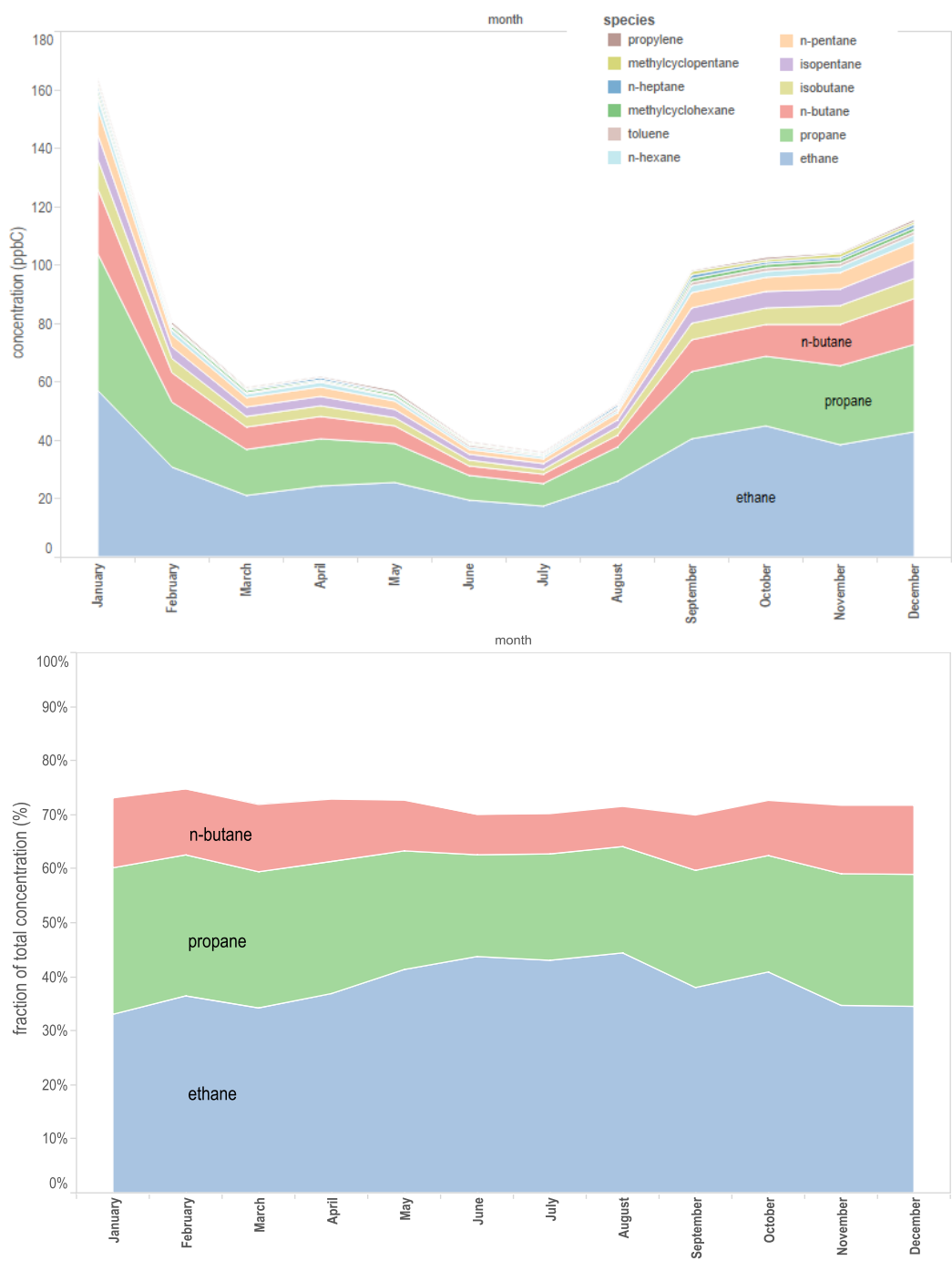


Figure 6-3. (a) Monthly average hydrocarbon concentrations at EML (ppbC); data are averaged over all hours and over 20 months of sampling. (b) Percentage of ppbC accounted for by ethane, propane and n-butane at EML

The concentrations show diurnal and monthly patterns, however, as shown in Figures 6-2b and 6-3b, the relative concentrations of the hydrocarbons are constant at all times of day and during all months, suggesting little seasonal or diurnal variability in emissions. The concentrations of light alkanes at EML are higher than at the other two sites. At EML the average morning maximum concentrations of ethane, propane, and butane total close to 100 ppbC, while at Flower Mound Shiloh and Hinton, the morning totals are 60 ppbC and 40 ppbC respectively. Summed over all days and hours in the 20 month sampling period, the average concentrations at EML for ethane, propane, and butane are 31.3, 19.4, and 9.2 ppbC, respectively. For Flower Mound Shiloh those averages are 18.4, 13.0, and 7.8, respectively. In the case of Hinton, the averages are 15.5 for ethane, 11.0 for propane, and 6.5 for butane.

The ethane to isopentane ratio, a measure of the relative importance of natural gas and gasoline related sources, is 7.54 at EML, while at Flower Mound Shiloh and Hinton, the ratio is 4.57, and 3.53, respectively. The difference in these ratios across the three sites highlights the predominant natural gas signal in the case of EML, and the higher impact of vehicle (gasoline) emissions in the urban (Hinton) and suburban (Flower Mound Shiloh) sites. The diurnal concentration patterns at the Flower Mound Shiloh and Hinton sites are shown in *Appendices, Section D.1*).

6.3.2 Background Corrected Alkane Concentrations at EML

In order to compare the observed hydrocarbon concentrations at EML to concentrations predicted from the Barnett Shale special inventory and a Lagrangian air quality model, it is necessary to correct the observed concentrations at EML for regional background concentrations. The ideal background site would exactly duplicate the distribution of natural emissions (e.g., geological seeps) and anthropogenic emissions not associated with natural gas production in the region surrounding the EML site. This hypothetical, ideal background site would also have meteorological conditions identical to the EML

site. Recognizing that all background sites will have limitations, two different background sites were used in this work. Data from both the Flower Mound Shiloh site and the Hinton site were used to estimate regional background concentrations, producing two sets of background corrected hydrocarbon concentrations.

The Flower Mound Shiloh site is only suitable as a background site when winds at the site are not from the Barnett Shale production region. Consequently, in estimating background concentrations using Flower Mound Shiloh observations, data were filtered for wind directions between 135° and 225° (53.4% of the total number of hours). This subset represents observations at Flower Mound Shiloh with the least influence from natural gas sources, mapped in Figure 6-1. Average diurnal hydrocarbon concentrations at Flower Mound Shiloh for observations with winds between 135° and 225° are provided in Appendices, Section D.3. For this data set, during the morning maximum, ethane, propane, and butane concentrations sum to less than 40 ppbC (similar to the observations at the Hinton site), compared to the 60 ppbC when all wind directions were considered. Taking this subset of observations, the average concentrations for ethane, propane, and butane are 13.3, 8.4, and 5.0, respectively, which are similar to the observations at the Hinton site.

No wind direction filtering was required when the Hinton site was used to establish background concentrations, since the location of the Hinton site is relatively far from the Barnett Shale production region. However, natural gas emission sources important in urban regions, such as leaks from natural gas distribution systems, may influence the Hinton data. The similarity of the alkane concentration data for the wind direction corrected Flower Mound Shiloh data set and the Hinton data set suggests that these urban sources are relatively small.

6.3.3 Comparisons of measurements and model predictions

Four different AERMOD scenarios were initially used to generate predicted VOC concentrations at EML. The scenarios considered sources included within different distances of the EML site (radii of 10, 25, 50, and 100 km) to assess whether predicted concentrations were dominated by sites near the measurement location. Table 6-2 shows the number of sources considered in each case, the total VOC emissions and the percentage relative to the total VOC reported in the Barnett Shale special inventory.

Table 6-2. Sources considered for each of the scenarios. The percentage shows the relationship between the input emissions for each case relative to the total emissions from all sources in the TCEQ inventory with known latitude and longitude.

Radius considered	Number of sites	Total emissions (tpy)	VOC Percentage of total VOC emissions
100 km	7,902	18,789	98.6%
50 km	5,168	11,221	58.9%
25 km	2,828	3,978	20.9%
10 km	670	430	2.26%

To be compared to AERMOD predictions, observations of hydrocarbon concentrations at the EML site were adjusted as follows.

- 1.) When Flower Mound Shiloh was used to determine background concentrations, data were filtered to consider only wind directions between 135° and 225° (53.4% of the original dataset, this subset represents observations at Flower Mound Shiloh with the least influence from natural gas sources). When Hinton was used as the background site, all wind directions were considered. Data from both Hinton and Flower Mound Shiloh were filtered to eliminate observations when wind speeds measured at EML were less than 0.2 m/s (hours with wind speeds equal or less than 0.2 m/s represent 0.3% of the original dataset). Dispersion models do not perform well at low wind speeds [10].
- 2.) All alkane concentrations in each hourly data record were summed, except for ethane (methane was not measured). The Barnett Shale special inventory included VOC estimates and the speciation profiles assign 83% of the emissions to alkanes, so it is expected that the reported VOC emissions are largely alkanes. Ethane was excluded because VOC emission inventories do not include ethane.
- 3.) Day and hour specific background concentrations for summed alkanes at the Flower Mound Shiloh site or the Hinton site were subtracted from the EML concentrations for the same day and hour; the cases where this difference was less than zero were filtered out for the analysis (2,552 hours, 26.5% of the original

dataset when Flower Mound Shiloh was used as a background site; 4,313 hours, 48.4% of the original dataset when Hinton was used as a background site). Cases where the concentration at EML is less than the concentration at the background are almost exclusively at the lower end of concentration distributions. The observations that are finally compared to AERMOD predictions represent 27.3% of the original dataset when Flower Mound Shiloh was used as a background site and 51.3% of the original dataset when Hinton is used as the background site.

Figure 6-4 shows a comparison between predicted VOC concentrations calculated using the AERMOD dispersion model and the observed, background corrected alkane concentrations as they were measured at the EML monitoring site.

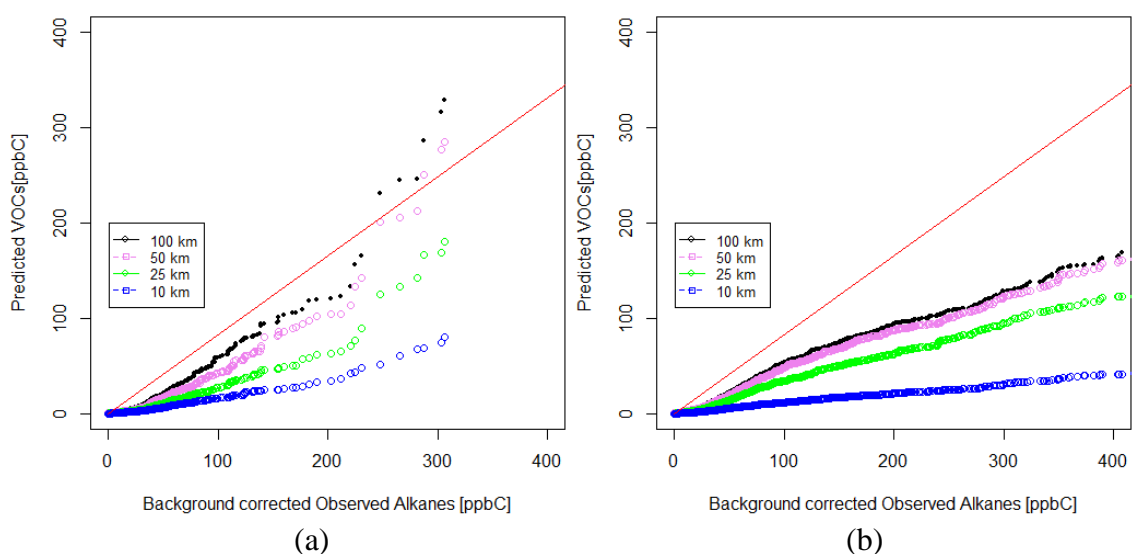


Figure 6-4. (a): Predicted total VOC concentrations vs. background corrected observed summed alkane concentrations at EML, when Flower Mound Shiloh was used as background site, for four sets of dispersion calculations, within 10, 25, 50, and 100 km radii of the observation site (b): Predicted total VOC concentrations vs. background corrected observed summed alkane concentrations at EML, when *Hinton was used as background site*, for four sets of dispersion calculations, within 10, 25, 50, and 100 km radii of the observation site. The red lines have a slope of 0.83 since 83% of the total VOC are speciated as alkanes.

Figure 6-4 shows that sources both close to and far from the EML site contribute to observed alkane concentrations at the site. The remainder of the analyses presented here will focus on analyses including all sources within a 100 km radius. The observations of total alkane concentrations are compared to 83% of the predicted concentrations of total VOC, since the speciation indicates that 83% of the emissions are paraffinic. As the radius of included sources is increased, the number of sources and the total emissions associated with the sources increase significantly (Table 6-2). Figure 6-4 shows considerable differences between the results as the radius is expanded, indicating that alkane concentrations at the EML site are not just reflecting emissions from sources in the near proximity of the EML monitoring station.

Differences between Figure 6-4a and 6-4b are driven by differences in the background data and in the sub-set of data that were used. For Figure 6-4a, the Flower Mound Shiloh site was used as a background and only data associated with predominantly southerly winds were used. As shown in Figure 6-1, the densest regions of natural gas production emissions are found north of the EML site; the subset of sources to the south of the site may not be representative of the sites to the north. In addition, the Flower Mound Shiloh site may be more heavily influenced by natural gas than the Hinton site, even when winds are from the south.

When using Flower Mound Shiloh as the background site, the regression of predicted vs. observed concentrations has a slope of 0.66 (standard error 0.013), for observations over 50 ppbC (background corrected), compared to an expected value of 0.83 (corresponding to an under-prediction bias of 21%). If the predicted values are increased by 10% to account for reported increases in natural gas production activity between 2009 and the 2010-2011 measurement period ²², the agreement between predicted and observed alkane concentrations shows even less difference.

When Hinton is used as background site, the agreement is not as good as the case of Flower Mound (Figure 6-5b); with a regression of predicted vs. observed concentrations

having a slope of 0.40 (standard error 0.005), for observations over 50 ppbC (background corrected) (corresponding to an under-prediction bias of 52%). If the predicted values are increased by 10% to account for reported increases in natural gas production activity between 2009 and the 2010-2011 measurement period, the slope of predicted versus observed concentrations is still only 0.43, relative to an expected value of 0.83. An additional analysis is included in Appendices (section D.3), where Hinton is used as background site, but data are filtered for the same hours that were considered in the final Flower Mound Shiloh dataset (which was filtered for wind directions). In this sensitivity analysis, the agreement with background corrected observations had a slope of 0.55, intermediate between the result considering all wind directions and the result using Flower Mound Shiloh, suggesting that some of the better performance of the inventory using Flower Mound Shiloh as a background is due to only considering sources to the south of EML.

Two additional sensitivity analyses were conducted to assess the potential contribution of under-estimated continuous emissions and episodic emissions to the predicted concentrations. For continuous emissions, results from a recent study of methane emissions from pneumatic controllers, chemical injection pumps, and equipment leaks in the Barnett Shale were used [8]. This study found that measured emissions from equipment leaks and pneumatic pumps and controllers were 159% higher in the Barnett Shale, compared to emissions estimated using current emission factors. Table 6-3 summarizes emissions per well derived from the measurements and compares these to emissions estimated based on a recent EPA greenhouse gas national inventory [11]. For estimated emissions of propane and higher alkanes, a weighted average gas composition for the Barnett Shale was used; 89.16% (vol.) methane, 4.24% ethane, and 2.67% C₃+ alkanes [12].

Table 6-3. Routine emissions from production sites in standard cubic feet (scf) of gas per minute per well.

	Pneumatic devices*	Chemical Injection Pumps*	Equipment Leaks*	Total	Total from EPA estimates**
Whole gas	0.569	0.152	0.071	0.792	0.202
Methane	0.507	0.136	0.064	0.707	0.180
Propane and higher alkanes	0.015	0.004	0.002	0.021	0.005

*Emission factors were obtained from supporting information of Allen, *et al*, and were based on activity counts from the same study[8].

**EPA estimates are based on values shown in Table 2 of Allen, *et al*. [8] Emissions from chemical pumps, pneumatic controllers, and equipment leaks, are 811 Gg of methane per year, or 80,365 scf/min of methane. Using the national gas well count (from the EPA national inventory) mentioned in Allen, *et al* of 447,000 wells; total methane emissions are 0.180 scf/min per well. Using the average gas composition for the Barnett Shale, the whole gas and C₃₊ emission rate estimates are derived (See Appendices, Section D.3).

Emissions of propane and higher alkanes, per well, from pneumatic devices, chemical injection pumps, and equipment leaks, are 0.021 scf/min, or 0.687 Mg/yr per well. Using the count of 14,886 producing wells[2], total propane and higher alkane emissions (VOC) from production sites would be anticipated to be roughly 10,200 tons per year. This is 159% higher than the corresponding estimate for similar categories in the TCEQ inventory of 3,935 tons per year (Table 6-1), and raises total VOC emissions by approximately 6,265 tons per year (33%). This change would eliminate the bias in the predicted concentrations when Flower Mound is used as a background site (causing an over-prediction bias of roughly 5%) and would reduce the bias to 37% if Hinton is used as a background site (all wind directions).

Episodic emissions due to events such as well unloadings are also not accounted for in the Barnett Shale emission inventory. Well unloadings are events that clear operating

wells from accumulated liquids that restrict the flow of natural gas. During the life of a well, as production declines, the velocity of natural gas in the well tubing might start to decrease, causing an accumulation of fluids in the wellbore. To remove the accumulated liquids, flow from the well is diverted into an atmospheric pressure tank, causing a large pressure drop that increases the velocity of the fluid. When there are no emission control systems in place, gas is vented to the atmosphere. Allen, *et al.* [8,12], report a total of 422 unloadings with venting per year for companies that operate a total of 2,791 wells in the Barnett Shale, a frequency of unloadings of 0.15 unloadings/well per year. Using the total number of wells reported by the TCEQ for the Barnett Shale in 2011 (14,886), the expected frequency of unloadings would be 2,251 events in a year.

To simulate the unloading events, 2,251 point sources from the TCEQ special inventory (within a 100 km radius from EML, from a total of the 8,025 inventoried sites) were randomly selected. Two unloadings scenarios are analyzed: (i) a frequency of unloadings resulting in 2,251 (random) point sources with 1 event with an emission per event of 1.1 Mg of methane (0.105 Mg propane and higher alkanes), and (ii) a frequency of unloadings resulting in 2,251 (random) point sources with 1 event with an emission per event of 3.7 Mg of methane (0.353 Mg propane and higher alkanes). Scenario (i) uses the average of emissions for an unloading event reported by Allen, *et al.*[8], while scenario (ii) uses the maximum observed emissions from an unloading event. Allen, *et al.* note that the unloading events sampled in their work may represent higher than average emissions, so the results reported here may overstate the impact of unloadings.

The time and date when each of the unloading events was simulated was randomly selected. Each event was assumed to last for one hour, and events were assumed to occur between 8 am and 5 pm, when operators would be onsite to perform manual unloading.

Figure 6-5 shows the propane and higher alkanes concentration at EML as a result of predicted unloadings. The plot shows the 1,597 distinct hours when the episodes occur (from the total 2,251 unloading events simulated, with a random date and time, some

events occur in the same day and hour, resulting in a total of 1,597 distinct hours where at least one event occurs).

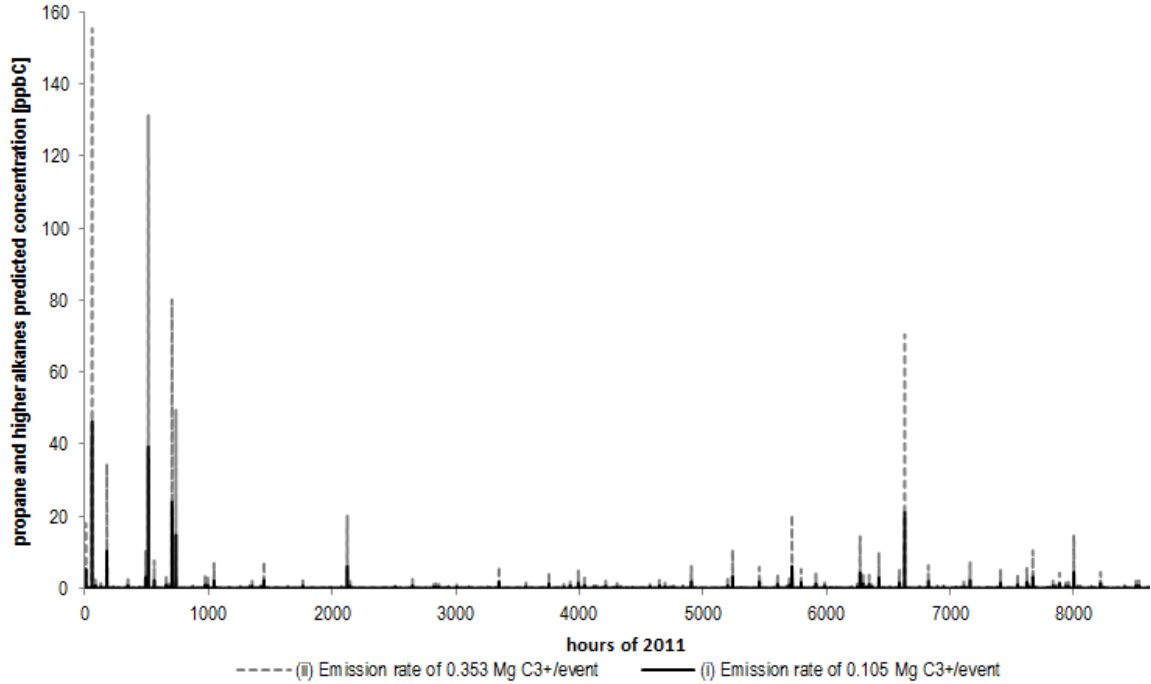


Figure 6-5. Time series showing the hours when an unloading event was simulated to occur and the resulting propane and higher alkanes predicted concentration at EML.

When the unloadings are simulated with the average emission rate of 0.105 Mg of propane and higher alkanes/event; the average C₃₊ predicted concentration at EML due to these events is 0.183 ppbC, with a median value of 0.007 ppbC and a maximum value of 46.4 ppbC, considering just the hours when an event is occurring (non-zero values only). When the maximum emission rate of 0.353 Mg propane and higher alkanes/event is used to simulate the unloadings, the average, median and maximum methane predicted concentration at EML is 0.613, 0.024, and 155 ppbC; respectively.

To compare these emissions to observations, the 4 largest simulated events are selected, with concentrations for scenario (i) of 21, 24, 39, and 46 ppbC, respectively. All the

events occurred under southeasterly/southwesterly winds (154° - 200°), with the exception of the second highest event which is related to northeasterly winds (4°). In all of these events, there are two sources with an unloading event during the same hour related to the predicted concentration at EML, with the exception of the event with a concentration of 39 ppbC; where there is just one source. All of the unloading were within a 30 km to 52 km radius of EML with the exception of one of the two sources related to the highest event (46 ppbC), which is 83 km away from EML. This analysis suggests that, given the assumptions stated above, if unloading events occur at the national average emission rate observed by Allen, *et al*, that several observations of increases in alkane concentrations greater than 20 ppbC should be evident in the dataset.

Close examination of Figure 6-4 shows that there are no episodic, anomalously low predictions from the dispersion models. In fact, when Flower Mound Shiloh is used as the background site, the dispersion model shows over-prediction at high concentration as opposed to the under-prediction at high concentration that would be expected if episodic events were important. If unloading events were occurring with an average emission of 0.105 Mg of propane and higher alkanes per event (scenario i) there should have been 4 events with 20 ppbC or larger enhancement of emissions. If scenario (ii) (higher emissions per event, 0.353 Mg of propane and higher alkanes per event) is considered, the number of events with a 20 ppbC or larger enhancement would be 8 over the sampling period. No such episodic anomalies are observed, indicating that unloadings or other episodic events are not significantly impacting ambient concentrations in the Barnett Shale. This result is specific to the Barnett Shale. In other regions, with higher frequencies of episodic events, events could play a role in the temporal distribution of ambient concentrations.

6.3.4 Benzene emissions

In addition to characterizing VOC emissions from natural gas production, the Barnett Shale Area Special Inventory reports emissions of benzene. A total of 54.5 tpy of benzene emissions (0.29% of the VOC emissions) are reported. Condensate tanks (44%),

water tanks (28%), and fugitives (20%) account for 92% of the benzene emissions. This distribution of sources by type is similar to VOC, where these source types were also among the top VOC emitters.

For the period of April 2010 to December 2011, benzene is not one of dominant species measured at EML. Table 6-4 shows the average, median, 90th percentile, 99th percentile, and maximum value from concentrations measured at EML and Flower Mound Shiloh and Hinton. Average, 90th and 99th percentile concentrations at Flower Mound Shiloh and Hinton are higher than at EML. When observed benzene concentrations at EML are compared to the predicted benzene concentrations (based on the AERMOD modeling) the estimated contribution from natural gas production ranges from a few percent of median total observed benzene concentrations to 30% of the total (99th percentile). This suggests that sources not directly associated with natural gas production determine benzene concentrations observed at EML. Some of these sources may be indirectly related to natural gas production, however, such as truck traffic servicing the production sites.

Table 6-4. Summary of benzene measured and predicted concentrations (ppbC) at the three sites.

Site	Average	Median	90th percentile	99th percentile	Maximum value
Eagle Mountain Lake	0.51	0.44	1.10	2.02	24.53
Eagle Mountain Lake PREDICTED from natural gas production	0.04	0.01	0.11	0.59	1.67
Flower Mound Shiloh	0.79	0.68	1.39	2.53	51.78
Hinton	0.94	0.71	1.79	4.64	13.84

6.4 Conclusions

In summary, hourly hydrocarbon concentrations from the Barnett Shale Natural Gas Production region indicate relatively little temporal variability in emissions from natural gas production. Hourly and daily variations in observed, background corrected concentrations were primarily explained by variability in meteorology, suggesting that episodic emission events, such as well unloadings, had little impact on hourly averaged concentrations. Total emissions for VOC are estimated to be approximately 25,300 tons/yr, when accounting for potential under-estimation of certain emission categories. This region produced, in 2011, approximately 5 bcf/d of natural gas (100 Gg/d or 40,000,000 tons/yr) for a VOC to natural gas production ratio (mass basis) of 0.00063. The dominant species are light alkanes.

6.5 References

- (1) Medlock, K. B. (2012) Modeling the implications of expanded US shale gas production. *Energy Strategy Reviews*, 1, 33-41.
- (2) Rail Road Commission of Texas, *Texas Gas Well Gas Production in the Newark, East (Barnett Shale) Field – 1993-2011*. Retrieved 9/11/2012 from http://www.rrc.state.tx.us/barnettshale/NewarkEastField_1993-2011.pdf
- (3) EIA, Monthly Natural Gas Gross Production Report; Retrieved 9/13/2012 from http://www.eia.gov/oil_gas/natural_gas/data_publications/eia914/eia914.html
- (4) Texas Commission on environmental Quality, *Barnett Shale Geological Area*; Retrieved 9/11/2012 from <http://www.tceq.texas.gov/airquality/barnettshale>
- (5) Texas Commission on environmental Quality, *Air Monitoring Sites*, Retrieved 9/11/2012 from http://www.tceq.texas.gov/airquality/monops/sites/mon_sites.html
- (6) Texas Commission on environmental Quality, *Point Source Emissions inventory*, Retrieved 9/19/2012 from <http://www.tceq.texas.gov/airquality/point-source-ei/psei.html#barnett2>
- (7) Texas Commission on environmental Quality, *Barnett Shale Special Inventory, Phase Two Workbook (Excel)*, Retrieved 10/7/2012 from <http://www.tceq.texas.gov/assets/public/implementation/air/ie/pseiforms/bshaleworkbook.xls>
- (8) Allen DT, Torres VM, Thomas J, Sullivan D, Harrison M, *et al.* (2013). Measurements of Methane Emissions at Natural Gas Production Sites, Proceedings of the National Academy of Sciences.
- (9) Perry, S.G.; Cimorelli, A. J.; Paine, R. J.; Brode, R W.; Weil, J. C.; Venkatram, A.; Wilson, R. B.; Lee, R. F.; R. F. (2005) AERMOD: A Dispersion Model for Industrial

Source Applications. Part II: Model Performance against 17 Field Study Databases; *Journal of Applied Meteorology*, 44, 682-693.

(10) Cimorelli, A. J.; Perry, S. G.; Venkatram, A.; Weil, J. C.; Paine, R. J.; Wilson, R. B.; Lee, R. F.; Peters, W. D.; Brode, R. W.; Paumier, J. O. (2004) AERMOD: Description of the Model Formulation, EPA, North Carolina.

(11) Environmental Protection Agency (EPA), Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2011. April 12, 2013 Retrieved 4/24/2013 from <http://www.epa.gov/climatechange/ghgemissions/usinventoryreport.html>

(12) Allen DT, Torres VM, Thomas J, Sullivan D, Harrison M, *et al.* (2013). Measurements of Methane Emissions at Natural Gas Production Sites: Study Appendices and Database, Austin, Texas: University of Texas <http://dept.ceer.utexas.edu/methane/study/>

7 DEVELOPING A METHANE, ETHANE, AND PROPANE EMISSION INVENTORY FOR THE BARNETT SHALE PRODUCTION REGION

7.1 Introduction

The natural gas supply chain is one of the largest anthropogenic sources of methane emissions in the United States, second only to enteric fermentation [1]. Due to the availability of new production technologies, such as horizontal drilling and hydraulic fracturing, U.S. natural gas production is increasing, making accurate estimates of methane emissions from natural gas production increasingly important.

Significant uncertainties exist in methane emission estimates; regional emission estimates based on ambient measurements are generally larger than regional emission estimates based on counts of emission sources and data on average, source specific, emission rates. Brandt, *et al.*[2] and Miller, *et al.* [3] have summarized recent emission estimates based on ambient measurements (top-down analyses) and conclude that emissions are generally higher than current inventories based on equipment counts and average emission factors (bottom-up estimates).

A challenge in reconciling top-down and bottom-up emission estimates is to identify which of many potential sources might be contributing to methane emissions inferred from top-down measurements. Since top-down measurements determine a regional total of methane emissions, contributions from a single source sector, such as the natural gas supply chain, are generally estimated by subtracting the contributions from all other sources (e.g., landfills, enteric fermentation) from the total regional estimate of emissions. This can introduce large uncertainties in emission estimates. An alternative is to use source specific tracers to estimate emissions from particular sources. Because natural gas is the only significant source of ethane and propane in the US, methane to ethane and methane to propane ratios have been used to attribute ambient methane emission observations to the natural gas industry [4-5]. Unfortunately, methane to ethane

and methane to propane ratios can vary spatially, even within a single natural gas field, making quantitative use of ratios difficult. Spatially resolved methane to ethane and methane to propane ratio data are needed to enable tracer analyses of natural gas emissions. [6-8].

This work provides spatially resolved methane, ethane, and propane emissions estimates as well as methane to ethane and methane to propane ratios for the Barnett Shale production region (North, Central Texas), scaled to the region's 2013 natural gas production levels. Development of spatially resolved methane, ethane and propane data for the Barnett Shale could improve source attribution based on airborne or ambient methane emissions measurements and inform comparisons between ambient measurements and bottom-up inventories.

This work estimates methane, ethane and propane emissions using a detailed inventory of VOC emissions prepared by the Texas Commission on Environmental Quality (TCEQ) and an extensive dataset of gas production data compiled as part of the direct measurement campaign conducted by Allen, *et al.* [9].

The Barnett Shale Special Inventory, prepared by the TCEQ, is a comprehensive inventory of Volatile Organic Compounds (VOC). It provides latitude, longitude, and magnitude of source – specific VOC emissions for over 8,000 sites [10]. By combining the 2009 Barnett Shale Special Inventory source locations, gas composition, and VOC emissions data with gas well composition data from a national dataset of methane emissions [9] from natural gas production, and scaling to 2013 production levels; this work develops estimates of methane, ethane, and propane emissions from over 18,000 individual sources in the Barnett Shale, spatially resolved on a 4km x 4km grid.

7.2 Methods

The Texas Commission on Environmental Quality (TCEQ) reported emissions from production sites in the Barnett Shale special inventory [10, 11] for the year 2009. This inventory compiles production data, source profiles, and emissions of volatile organic compounds (VOC). VOCs include propane and heavier hydrocarbons (methane, ethane are not inventoried because by regulatory definition, they are not VOCs). VOC data were collected for 18,466 sources; accounting for a total of 19,050 tons per year (tpy) (92% of the emissions are source-specific, with the remaining 8% being site-specific only). The source specific data are aggregated onto 8,025 upstream sites (with geospatial information).

Emissions are divided into 16 source categories, with condensate tanks, fugitives, water tanks, and engines dominating the VOC emissions. Figure 7-1 shows the distribution of VOC emissions by source category as reported in the special inventory.

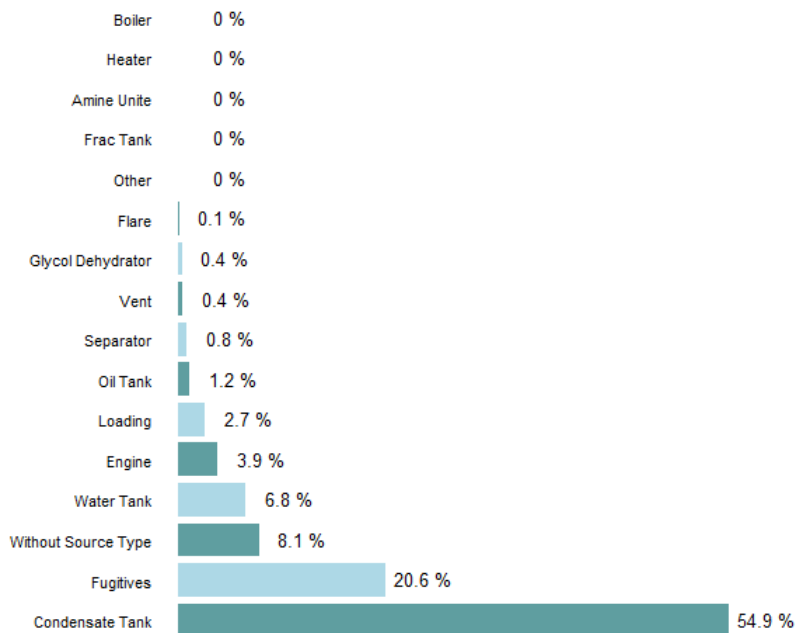


Figure 7-1. VOC emissions by source type, as reported in the TCEQ special inventory. Total emissions add up to 19,050 tpy.

For the development of the inventory, the TCEQ provided an emission calculator, allowing for estimates of emissions using site-specific data (summing up emissions from sources contained within each site) [12]. On-site direct measurements were preferred to indirect measurements or emission-factor based methods but they were not required [13]. The inventory of methane, ethane and propane emissions developed in this work uses the TCEQ special inventory (for VOC emissions) together with natural gas composition data to estimate methane, ethane and propane emissions.

Allen, *et al.* [9,14] reported gas composition for 150 wells in 4 different production regions across the U.S. Figure 7-2 shows the linear regression of percent of methane in the produced gas (% C1) as a function of percent VOC in the produced gas (% VOC, for propane and heavier hydrocarbons), ($slope = -1.978$, $intercept = 96.149$, $R^2 = 0.962$).

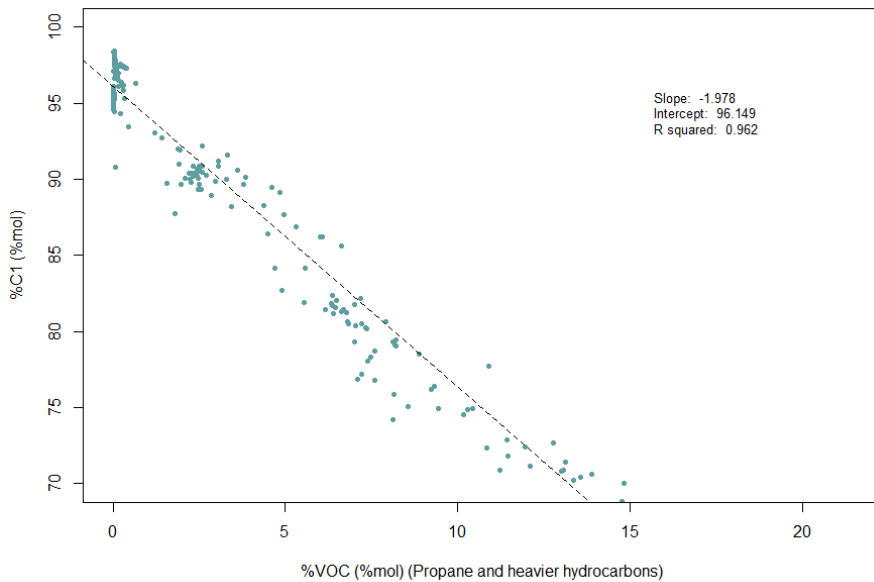


Figure 7-2. Linear regression of %C1 as a function of %VOC for 150 wells across the U.S. where direct measurements were made by Allen, *et al.* [14].

The linear relationship observed for wells in different regions allows the estimation of the percent of methane in vented gas if the percent of VOC is known, regardless of the wet or dry nature of the well and the specific regional characteristics. Similarly, the linear

regression of percent propane in gas (%C3) as a function of %VOC shows that roughly 60% of the VOC is propane (*slope* = 0.588, *intercept* = -0.046, $R^2 = 0.983$). Thus, if the %VOC for a site is known, it is possible to estimate %C1 and %C3 from the regressions presented in this work, and consequently estimate %C2 from the difference (100% - [%C1 + %VOC]).

As part of the data requested for the TCEQ special inventory %VOC values were collected for sites that reported fugitive emissions. Only 30% of the roughly 8,000 sites report %VOC value; nonetheless, the sites with known %VOC have a characteristic spatial distribution throughout the Barnett Shale region, therefore, missing %VOC values were spatially interpolated using an inverse distance weight method (IDW). (See Appendices, *Section E.1* for more details on the interpolation of percent VOC values).

Using the interpolated %VOC values, and the correlations presented in this work, %C1, %C2, %C3 were estimated to determine the composition of the gas at each site. Table 7-1 summarizes how gas composition, source-specific VOC emissions, and site-specific production data were used to estimate methane, ethane, and propane emissions, following a specific rationale based on the specific characteristics of each specific emission source category. Appendices (Section E.2) describes in detail how each separate source category was treated). All emissions were estimated for the baseline year, 2009, and then projected to 2013.

Table 7-1. Source-specific rationale for the estimation of methane, ethane, and propane, emissions from each source category. Section E.2 from the Appendices explains each calculation in detail.

Source Category	Reported VOC Emissions (tpy)	Rationale for the estimation of Methane emissions
Condensate Tank	10,465	Estimated the solubility of gas in oil (Vasquez-Beggs equation [14, 15] and used site-specific production data to determine the volume of gas that flashes. It was assumed that all methane, ethane and propane flash, and that their relative composition is similar to their relative composition in the produced gas.
Fugitives	3,935	Corrected emissions for the underestimation of equipment leaks and pneumatic devices [16], and then methane emissions were scaled based on % VOC.
Water Tank	1,286	Estimated the solubility of methane, ethane, and propane in water (Henry's law [14]), and used site-specific information production data to determine the volume of gas vented from water tanks.
Engine	749	Methane, ethane, and propane emissions were scaled based on % VOC and reported VOC emissions.
Loading	511	Assumed no methane emissions are released from these source category.
Oil Tank	221	Estimated the solubility of gas in oil (Vasquez-Beggs equation [14, 15]) and used site-specific production data to determine the volume of gas that flashes. It was assumed that only methane, ethane and propane flash, and that their relative composition is similar to their relative composition in the produced gas.
Separator	161	Methane, ethane, and propane emissions were scaled based on % VOC and reported VOC emissions.
Vent	84	Methane, ethane, and propane emissions were scaled based on % VOC and reported VOC emissions.

Table 7-1. (continued).

Source Category	Reported VOC Emissions (tpy)	Rationale for the estimation of Methane emissions
Glycol Dehydrator	66	Methane, ethane, and propane emissions were scaled based on %VOC and reported VOC emissions.
Flare	20	Methane, ethane, and propane emissions were scaled based on %VOC and reported VOC emissions.
Other	9.5	Methane, ethane, and propane emissions were scaled based on %VOC and reported VOC emissions.
Frac Tank	3.5	Emissions were estimated based on the estimated emissions from completion flowback events measured by Allen, <i>et al.</i> [9, 15]
Amine Unite	2.5	Methane, ethane, and propane emissions were scaled based on %VOC and reported VOC emissions.
Heater	1.7	Methane, ethane, and propane emissions were scaled based on %VOC and reported VOC emissions.
Boiler	0.30	Methane, ethane, and propane emissions were scaled based on %VOC and reported VOC emissions.
Without Source Type	1,535	Methane, ethane, and propane emissions were scaled based on %VOC and reported VOC emissions..

7.3 Results and Discussion

7.3.1 Spatially resolved emission inventory

To facilitate the use of the inventory developed in the present work as a tool to reconcile top-down and bottom-up measurements, emissions were summed up and divided into 4 km by 4 km grid cells that correspond to the grid cells used in TCEQ air quality modeling applications, which were developed for the State Implementation Plan (SIP) in the Dallas-Fort Worth ozone nonattainment area [17].

Total methane emissions for the Barnett Shale production region were estimated to be 5,741 kg/h (2,619 MMscf/yr) for 2009. Roughly 60% of the emissions are attributed to fugitives, with 12% from engines, 7% from condensate tanks, 6% from water tanks and 13% without source type. Figure 7-3 illustrates the spatial distribution of methane emissions; plotted as density of emissions (kg/h per square kilometer) (Tables with estimated emissions for each grid cell are provided in Appendices, Section E.3).

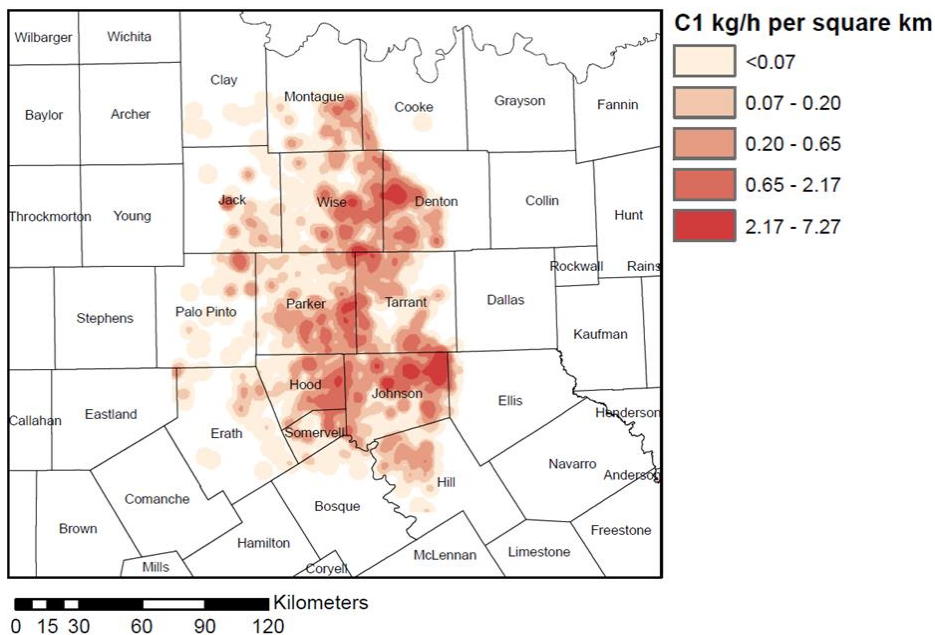


Figure 7-3. Density of estimated methane emissions (2009) (kg/h per square kilometer).

The spatial distribution of the estimated methane emissions indicates two hotspots: one on the Wise\Denton county border, and the other in the northeast corner of Johnson County. The hotspot in Wise and Denton counties located in the region of the Barnett Shale with the highest well density and with the highest VOC emissions [12]. On the other hand, Johnson County has a lower density of wells, hence the methane hotspot is explained by the combination of significant fugitive emissions combined with a dry gas composition. Figure 7-4 shows the distribution of VOC fugitive emissions and %VOC by county. The distribution of VOC emissions for Johnson County is similar to other counties, but the distribution of %VOC for sites in Johnson county is characterized by lower values than the rest of the region (gas in Johnson is drier). Thus for a given amount of VOC emissions, whole gas emissions would be higher in Johnson County and consequently, methane emissions would also be scaled towards higher values.

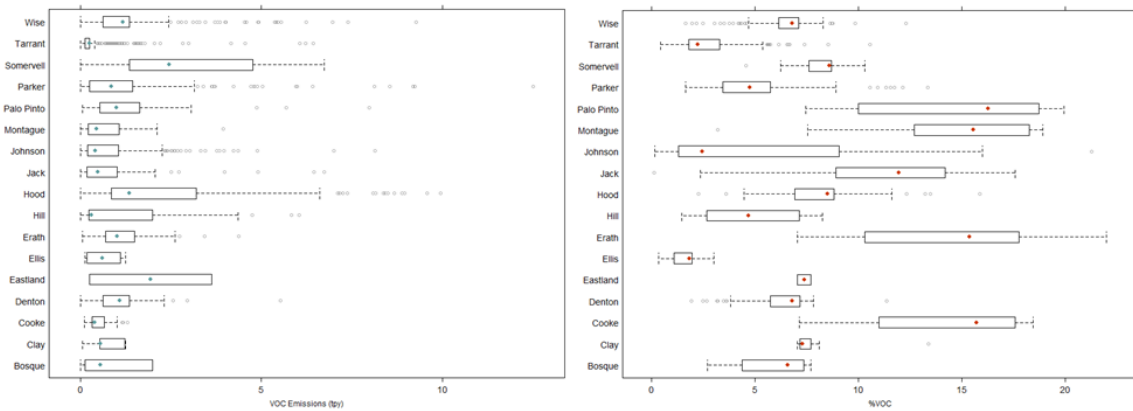


Figure 7-4. Distribution of VOC emissions (*left*) and %VOC (*right*) by county. The distributions are represented by boxplots, with the central dot representing the median value, and the range of the rectangle representing the interquartile range (25th and 75th percentile).

7.3.2 Scaling emissions

To scale emissions from the baseline year (2009) to 2013, it was assumed that, since emissions are associated with wells in routine operation, emissions can be scaled based on natural gas production. Natural gas production for the Barnett Shale in 2009 was 4,921

MMcf/d, while for 2013, production was 5,309 MMcf/d [11], an 8% increase in production relative to 2009.

The scaled 2013 methane emissions are equal to the product of an emission factor (EF); defined as the ratio of methane emissions for the baseline year, divided by gas production for the baseline year, and an activity factor (AF), in this case; natural gas production for 2013.

If an average emission factor is calculated as the sum of all estimated methane emissions in the domain (2,619 MMscf/yr), divided by 2009 Natural Gas Production from all the sites in the domain (1,790,655 MMscf/yr), the EF would be 0.0015, which multiplied by the activity factor of 1,937,785 MMscf/yr (Natural Gas Production 2013) would yield total 2013 methane emissions of 2,834.6 MMscf/yr or 6,213 kg/h.

Figure 7-5 shows the distribution of emission factors for each grid cell;. 12% of the grid cells had an emission factor of zero. For non-zero EF grid cells, the median value was 0.0022, the average value was 0.0154, with a 25th percentile of 0.0006, a 75th percentile of 0.006, and a 95th percentile of 0.0522.

57% of the 2013 Natural Gas Production is in cells with an emission factor less than 0.001. For the remaining Natural Gas Production, 35% is in cells with an emission factor between 0.001 and 0.01, and 8% has an emission factor greater than 0.01. Although more than half of the production falls into cells with an emission factor that is below the median, the skewed distribution of emission factors causes the estimated emissions to depend strongly on the production in the cells with the highest EFs. (See Section E.3 of Appendices for tables reporting EF and AF values for each grid cell).

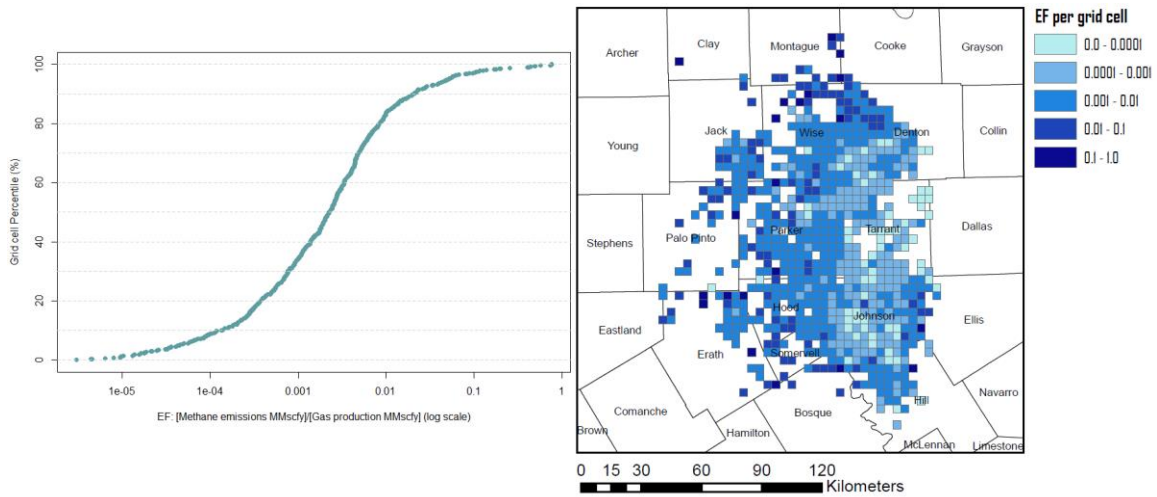


Figure 7-5. Cumulative distribution plot for non-zero emission factors for each grid cell (log scale) (*left*), and map showing the spatial distribution of emission factors in the Barnett Shale (*right*).

Since estimated emissions have a strong correlation with natural gas production, and the skewed distribution of emission factor evidences important regional differences, the relationship of EF and AF for each grid cell is important. Therefore, another way to estimate the scaled emissions, is to obtain the product of EF and AF for each individual grid cell and then sum all the emissions. Following this method, estimated methane emissions for 2013 are 46,745 kg/h (compared to 6,213 kg/h when regional averages were used).

In summary, the Barnett Shale exhibits a heterogeneous distribution of methane emissions. The region's heterogeneity is driven by both the skewed spatial distribution of emission factors among grid cells and the heterogeneous distribution of the temporal changes in production from 2009 to 2013. Appendices, Section E.4 analyses the spatial distribution of both emission factors and activity factors. These significant differences within the Barnett Shale production region should be taken into account when top-down and bottom-up measurements are compared.

Using methods similar to those used for methane, emissions for ethane and propane were estimated. 2009 emissions for ethane were 778 kg/h and for propane were 599 kg/h. 2013 projected emissions were 2,848 kg/h for ethane and 2,301 kg/h for propane (if emissions are scaled for each individual grid cell). As mentioned previously, methane to ethane and methane to propane ratios have been used extensively to attribute ambient methane emission observations to the natural gas industry. Figure 7-6 and Figure 7-7 show the spatially resolved methane to ethane and methane to propane ratios for the Barnett Shale, respectively. (Data set with ethane and propane emissions, as well as ratios is provided in Appendices, Section E.3).

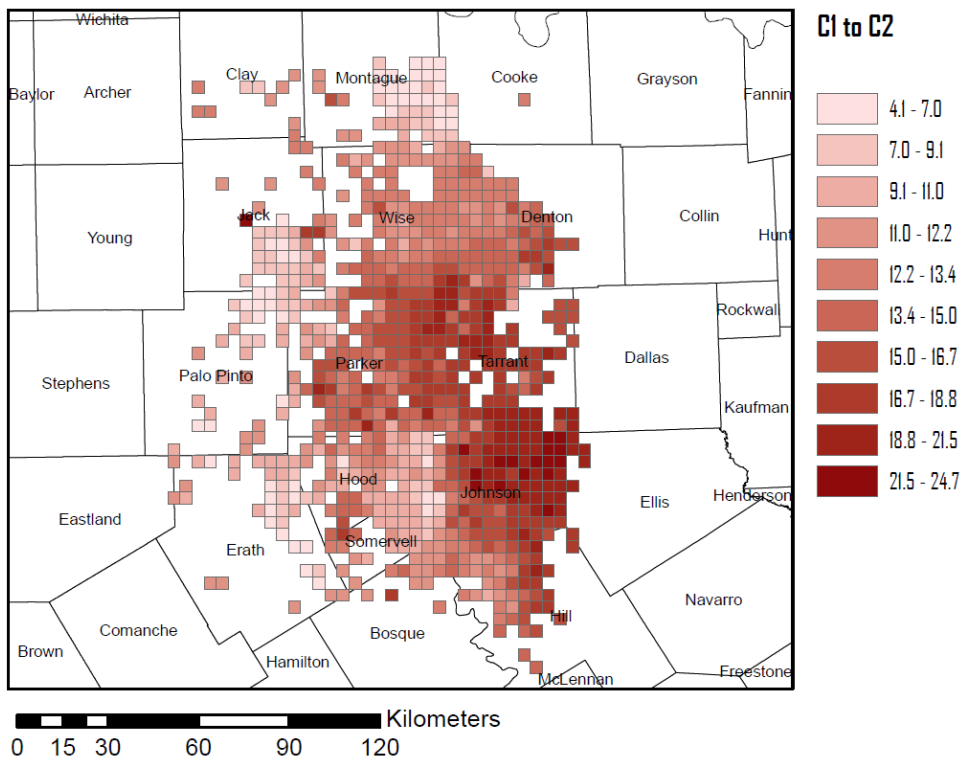


Figure 7-6. Methane to ethane ratio of emissions.

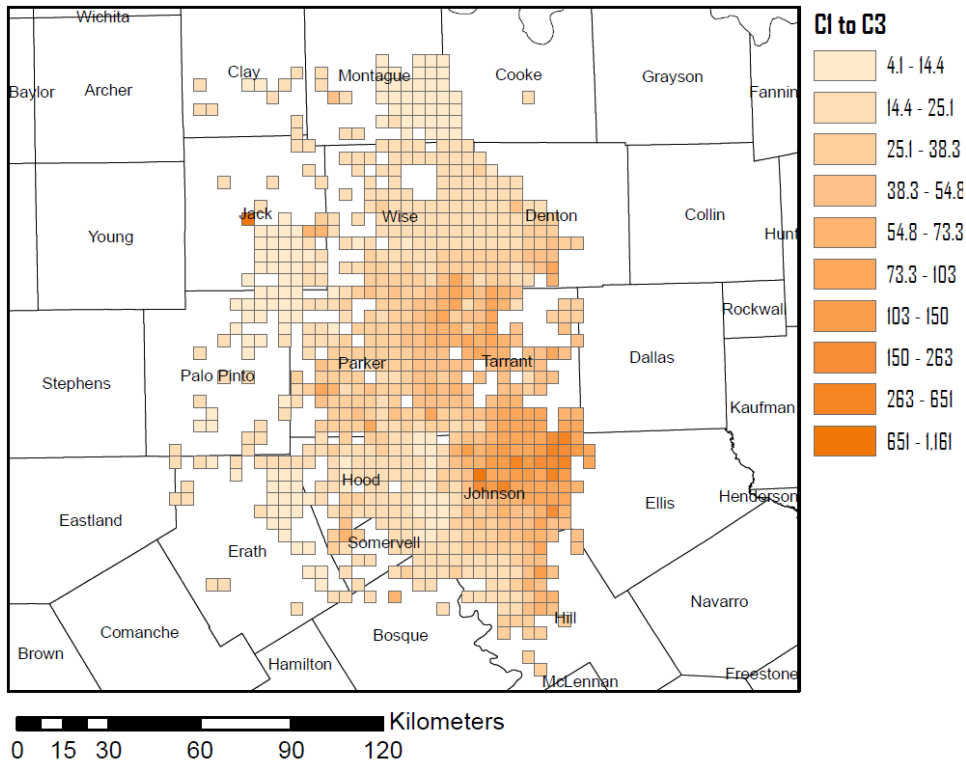


Figure 7-7. Methane to propane ratio of emissions.

7.4 Conclusions

Methane, ethane, and propane emission inventories can be used for the reconciliation of bottom-up and top-down measurements in the Barnett Shale Production Region. Significant shifts in the spatial distributions of production should be taken into account when direct and indirect measurements are compared against estimates from emission inventories.

The development of spatially resolved methane, ethane and propane data for the Barnett Shale could have particular relevance to a diverse group of research teams that are utilizing airborne, vehicle, and ground-based measurements to estimate emissions in the Barnett Shale.

7.5 References

- (1) Environmental Protection Agency (EPA). Inventory of U.S. Greenhouse Gas Emissions and Sinks. <http://www.epa.gov/climatechange/Downloads/ghgemissions/US-GHG-Inventory-2014-Main-Text.pdf> (accessed 21 June 2014).
- (2) Brandt AR, Heath GA, Kort EA, O’Sullivan F, Pétron G, Jordaan, SM, Tans P, Wilcox J, Gopstein, AM, Arent D, *et al.*(2013) Methane Leaks from North American Natural Gas Systems. *Science*, 343: 733-735.
- (3) Miller SM, Wofsy SC, Michalak AM, Kort EA, Andrews AE, Biraud SC, Dlugokencky EJ, Eluszkiewicz J, Fischer ML, Janssens-Maenhout G, *et al.*(2013) Anthropogenic emissions of methane in the United States. *Proc. Natl. Acad. Sci. U.S.A.*, 110, 20018–20022.
- (4) Yaping, X.; Logan, J.; Jacob, D.; Hudman, R.; Yantosca, R.; Blake, D. Global budget of ethane and regional constraints on U.S. sources. *J. Geophys Res.* **2008**, 113, D21306.
- (5) Aydin, M.; Verhulst, K. R.; Saltzman, E. S.; Battle, M. O.; Montzka, S. A.; Blake, D. R.; Tang, Q.; Prather, M. J. Recent decreases in fossil-fuel emissions of ethane and methane derived from firm air. *Nature* **2011**, 476 (7359), 198-201.
- (6) Miller, S.; Wofsy, S.; Michalak, A.; Kort, E.; Andrews, A.; Biraud, S.; Dlugokencky, E.; Eluszkiewicz, J.; Fischer, M.; Janssens-Maenhout, G.; Miller, B.; Miller, J.; Montzka, S.; Nehrkorn, T.; Sweeney, C. Anthropogenic Emissions of Methane in the United States. *Proc. Natl. Acad. Sci. U.S.A.* 2013, 110 (50), 2018-2002.
- (7) Kort, E. A.; Eluszkiewicz, J.; Stephens, B.; Miller, J.; Gerbig, C.; Nehrkorn, T.; Daube, B.; Kaplan, J.; Houweling, S.; Wofsy, S. Emissions of CH₄ and N₂O over the United States and Canada based on a receptor-oriented modeling framework and COBRA-NA atmospheric observations. *Geophys. Res. Lett.* 2008, 35, L18808.
- (8) Pétron, G.; Karion, A.; Sweeney, C.; Miller, B.; Montzka, S.; Frost, G.; Trainer, M.; Tans, P.; Andrews, A.; Kofler, J.; Helmig, D.; Guenther, D.; Dlugokencky, E.; Lang, P.;

Newberger, T.; Wolter, S.; Hall, B.; Novelli, P.; Brewer, A.; Conley, S.; Hardesty, M.; Banta, R.; White, A.; Noone, D.; Wolfe, D.; Schnell, R. A new look at methane and nonmethane hydrocarbon emissions from oil and natural gas operations in the Colorado Denver –Julesburg Basin. *J. Geophys. Res. Atmos.* **2014**, *119*, 1-17.

(9) Allen, T. D., Torres, M. V., Sullivan, D., Harrison, M., Hendler, A., Herndon, S. C., et al. (2013). Measurements of Methane Emissions at Natural Gas Production Sites. *Proceedings of the National Academy of Sciences*. 110, 17768-17773.

(10) Texas Commission on Environmental Quality (TCEQ). Barnett Shale Special Emissions Inventory Phase I. 2011. <https://www.tceq.texas.gov/assets/public/implementation/air/ie/pseiforms/Barnett%20Shale%20Area%20Special%20Inventory.pdf> (accessed 21 June 2014).

(11) Texas Commission on environmental Quality, Barnett Shale Geological Area; Available at <http://www.tceq.texas.gov/airquality/barnettshale> (accessed 9/11/2012)

(12) Texas Commission on environmental Quality, Barnett Shale Special Inventory, Phase Two Workbook (Excel), Available at <http://www.tceq.texas.gov/assets/public/implementation/air/ie/pseiforms/bshaleworkbook.xls> (accessed 10/7/2012).

(13) Texas Commission on environmental Quality, Point Source Emissions inventory, Available at <http://www.tceq.texas.gov/airquality/point-source-ei/psei.html#barnett2> (accessed 9/19/2012).

(14) Allen DT, Torres VM, Thomas J, Sullivan D, Harrison M, et al. 2013. Measurements of Methane Emissions at Natural Gas Production Sites: Study Appendices and Database. Austin: Univ. Tex. <http://dept.ceer.utexas.edu/methane/study/>

(15) Texas Commission on Environmental Quality, Calculating Volatile Organic Compounds (VOC) Flash Emissions from Crude Oil and Condensate Tanks at Oil and Gas Production Sites, Available at

http://www.tceq.texas.gov/assets/public/permitting/air/Guidance/NewSourceReview/guidance_flashemission.pdf (Accessed 6/22/2014).

(16) Zavala-Araiza, D., Sullivan, D., & Allen, D. T. (2014). Atmospheric hydrocarbon emissions and concentrations in the Barnett Shale natural gas production region. *Environmental Science & Technology*, 48(9), 5314-5321.

(17) Texas Commission on environmental Quality, Dallas-Fort Worth Eight-Hour Ozone SIP Modeling (2006 Episode), Available at <http://www.tceq.state.tx.us/airquality/airmod/data/dfw8h2> (accessed 6/22/2014)

8 DESIGN OF A MULTI-SCALE METHANE EMISSION DETECTION AND ATTRIBUTION IN NATURAL GAS PRODUCTION REGIONS

8.1 Introduction

As regulations are promulgated to reduce methane emissions from natural gas production sites [1-4], it becomes important to track the effectiveness of the emission reduction programs. As documented in previous chapters, methane emissions from natural gas production occur on a variety of spatial and temporal scales [5, 6], and so systems designed to monitor emission reductions must be capable of detecting changes in both emissions from routine operations (e.g. equipment leaks, pneumatic devices) and from episodic events (e.g. well completions, liquids unloading). Figure 8-1 simulates a representative emission signal from a natural gas production well pad where: (i) there is an equipment leak with an average whole gas emission rate of 10scf/h, (ii) an intermittent-vent controller with an average whole gas emission rate of 28 scf/h, and (iii) a manual liquid unloading event takes place with a duration of roughly 1.2 hours, and a total vented volume of gas of 120,000 scf. Figure 8-2 highlights, on different scales, the equipment leak signal (*left*), and the liquid unloading event signal (*right*).

This chapter describes a potential monitoring system, with three spatial tiers, for identifying methane emission sources in natural gas production regions, such as those illustrated in Figures 8-1 and 8-2. The design of a possible methane monitoring network for the Barnett Shale region will be used as an illustrative case study. A network of high sensitivity, high time resolution fixed ground monitors could be used to identify large intermittent emissions, and could be coupled with a mobile unit that will allow a high sensitivity identification and quantification of specific emission sources.

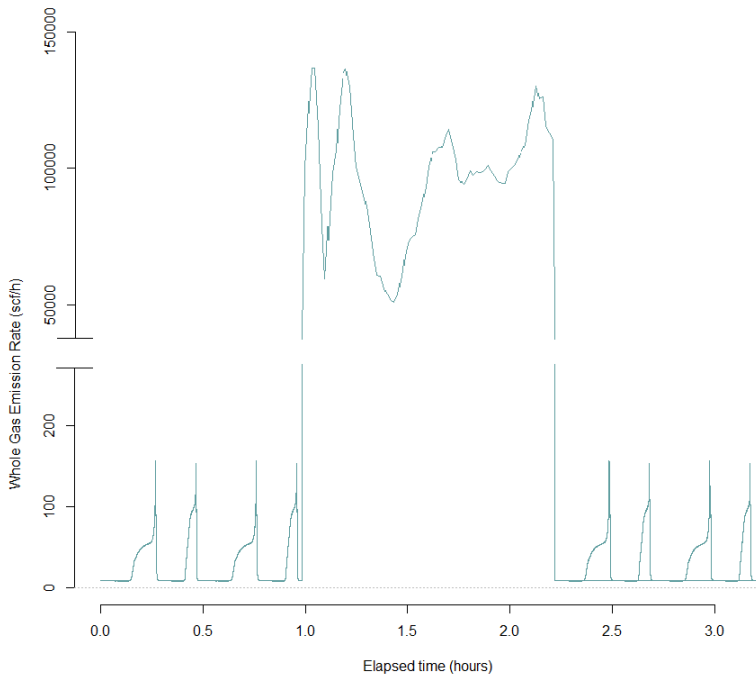


Figure 8-1. Simulated emission signal from a natural gas well pad. Methane emissions caused by different sources are illustrated. Actuations from an intermittent-vent pneumatic controller occur as part of the routine operation of the well. The offset between the actuations of the pneumatic controller and the zero emissions baseline is caused by the existence of an equipment leak on the site. At 1.0 hours (elapsed time) a manual liquid unloading event takes place, with duration of 1.2 hours and a 120,000 scf of whole gas emitted. After the event ends, the well goes back to routine operation, the equipment leak and the emissions from the pneumatic device and leak are apparent again.

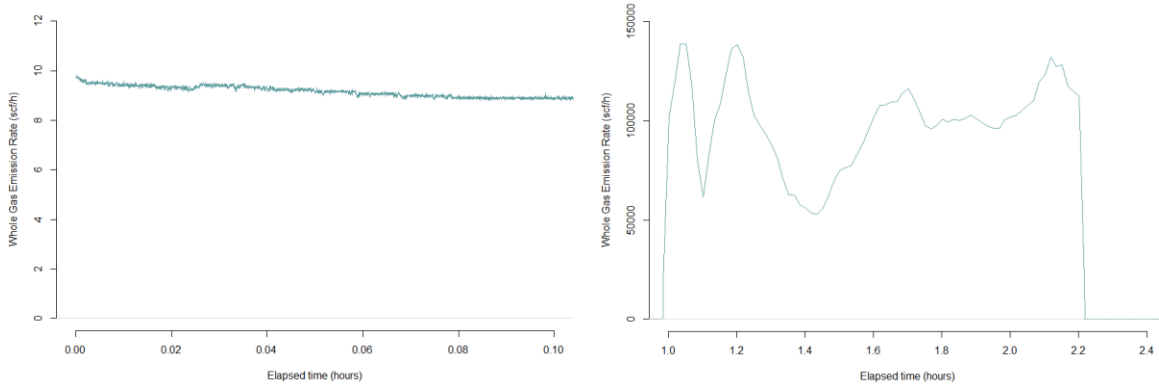


Figure 8-2. Leak with an average whole gas emission rate of 10 scf/h (left), and liquid unloading event, with total emissions of 120,000 scf for the event.

8.2 Design of the monitoring system

8.2.1 Study Region

The Barnett Shale production region has shown a significant growth in gas production activity during the last ten years; with production of 0.11 billion cubic feet per day in 2000, increasing to 5 billion cubic feet per day by 2011 [7, 8]. Total natural gas withdrawals in the US in mid-2011 were approximately 70 billion scf/d (bcf/d). With over 10,000 well sites in a mature production region, the Barnett Shale was selected as a potential location for a proof of concept regional methane monitoring network. *Chapter 6* of this thesis presented analyses on VOC emissions from this region, and *Chapter 7* expanded those analyses by developing a methane inventory; which is used in the this chapter to select the specific section of the Barnett Shale where a proof of concept network could be placed.

Figure 8-3 shows well density and methane emissions density for the Barnett Shale region. Based on the locations of methane emission hotspots, a study region covering 1,000 – 2,000 wells was selected (black circle in Figure 8-3, expanded in Figure 8-4). The characteristics of the monitoring network shown in Figure 8-4 are described in the next section.

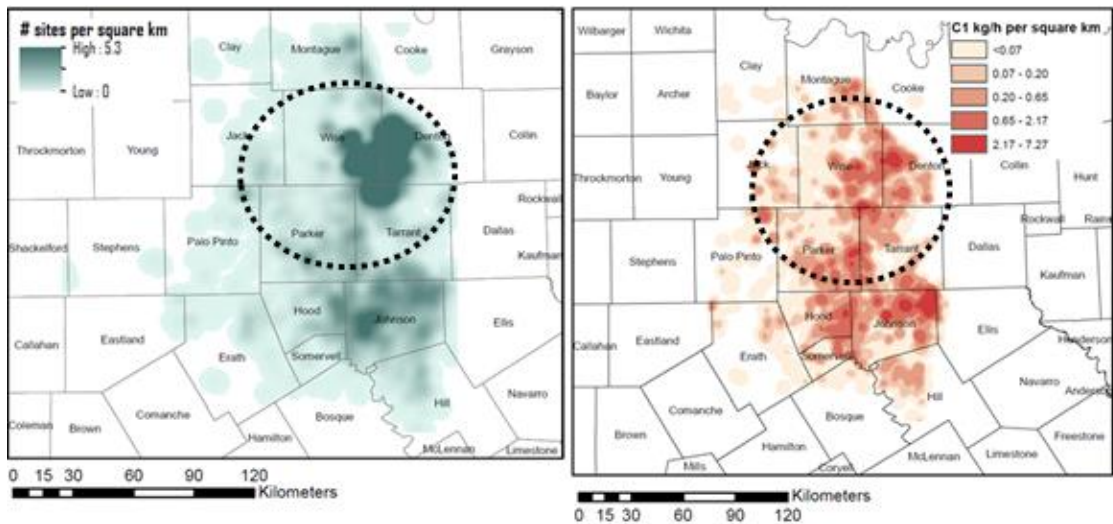


Figure 8-3. Well density (*left*) and Methane emissions density (*right*) for the Barnett Shale production region. Methane emissions were estimated following the procedure described in *Chapter 7* of the present work. The black circle shows the study region (1,000 – 2,000 wells).

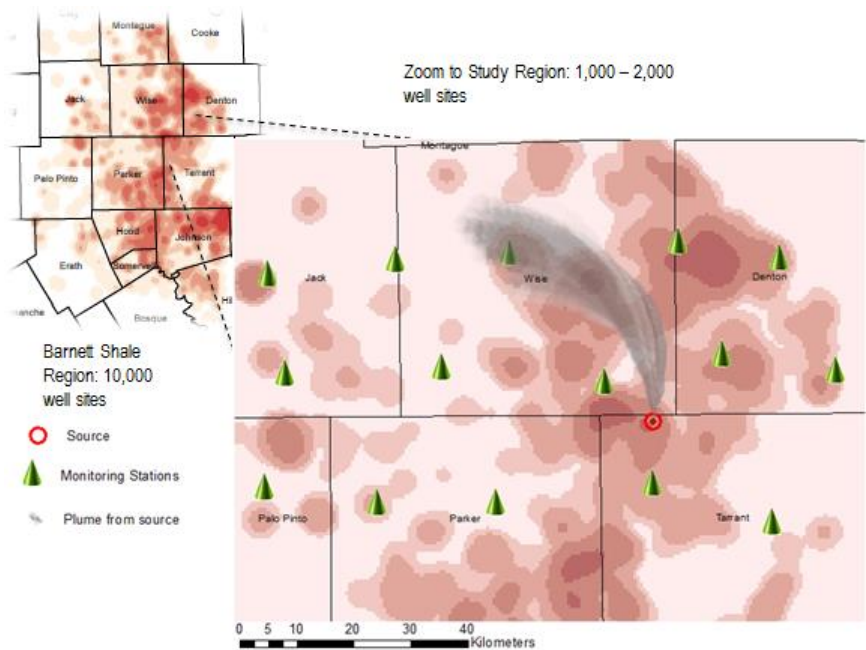


Figure 8-4. Spatial location of the study region, with monitoring network that detects methane hotspots (density of methane emissions in the Barnett Shale region is shown in red shading).

8.2.2 *First level: Monitoring network*

At the largest spatial scale, fixed ground monitors could be deployed at multiple sites in production basins, as shown in Figure 8-4, to identify changes in emissions. Methane and ethane could be measured at 1 Hz, using Quantum Cascade Tunable Infrared Laser Differential Absorption Spectroscopy (QC-TILDAS); wind speed, wind direction, and carbon dioxide could also be measured using current measurement technologies.

The QC-TILDAS system is an atmospheric trace gas detector based on mid-infrared absorption spectroscopy that uses pulsed quantum cascade lasers. Tunable infrared laser differential absorption spectroscopy systems (TILDAS) are characterized by a high sensitivity, high specificity and fast response time in the mid-infrared spectral region. By incorporating the use of quantum cascade lasers, it is possible to operate the system at near room temperature. The system has been used for detection of nitric oxide [9], carbon monoxide, and methane [10], among other atmospheric trace species.

These instruments employ continuous, quantitative spectroscopic ethane and methane measurements in the fingerprint mid-IR with extreme precision and speed; the 1-second detection sensitivities are 400 and 150 parts per trillion for methane and ethane respectively [11].

8.2.3 *Second level: Mobile Unit*

At intermediate spatial scales, vehicle mounted measurements could be used to quantify emissions at specific sites; the site emission total could then be compared to detailed, site specific emission inventories as described in *Chapter 7*. The vehicle mounted monitors could include QC-TILDAS system for methane and ethane, as well as wind speed and wind direction data, operating at 1 Hz.

Figure 8-5 illustrates the role of the mobile unit in detecting emissions from specific well sites.

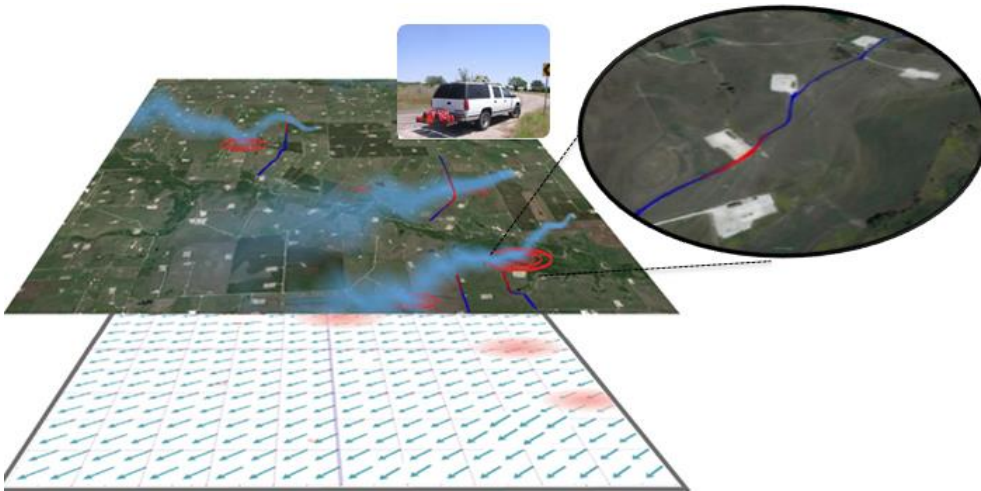


Figure 8-5. Mobile unit detects emissions at specific sites. Driving path shaded accordingly to the magnitude of emissions.

8.2.4 Third level: IR Camera and conventional leak measurement methods

At the finest spatial scales, two approaches could be used. For sites where a vehicle mounted system has quantified emissions higher than expected for a site, infrared camera systems could be used to identify the precise location of the leak; where possible, conventional leak measurement methods (e.g., HiFlow © measurements) will be used to confirm the emission rate determined by the vehicle mounted system.

As described in Chapter 3 of this thesis and in Allen, *et al.* [5], the Hi Flow® Sampler is a portable, intrinsically safe, battery-powered instrument that has been used for several decades in measuring emissions of methane in the natural gas supply chain [12-14].

Figure 8-6 shows examples of the IR camera detecting specific emission sources at a well site.



Figure 8-6. IR camera pinpoints specific emission source. Examples show equipment leak (*left*), venting from tank (*center*), and emissions from pneumatic controller (*right*).

8.2.5 *Coupling the different monitoring levels*

Instantaneous measurements of methane and ethane concentrations at fixed sites could be continuously compared to predicted concentrations. Predicted concentrations generated by lagrangian models could use, as input, site specific emission data for thousands of well sites in a production region and instantaneous wind field measurements generated by the network of fixed ground sites. *Chapter 6* of the present work describes a similar approach, where VOC concentrations at fixed receptor site were compared against predicted concentrations generated by a langrangian dispersion model. In this case, the methodology could be applied to the network of sites and in a continuous manner. Each hour, back trajectories could be calculated for each fixed ground site based on the local wind fields measured at the sites. Methane and ethane concentrations, as well as concentrations of key tracer species such as CO₂ (for combustion sources), could be predicted every minute, based on the hourly back trajectories and detailed emission inventories, and compared to observations. Daily routes for the vehicle mounted sensors could be mapped based on these daily identifications of source regions yielding higher than expected concentrations.

8.3 Conclusions

Monitoring methane emissions from natural gas production is a complex task due to the heterogeneous spatial and temporal patterns of the emission sources. This Chapter has described a set of monitoring and attribution systems at different scales, allowing the interpretation, characterization, and accurate detection of the emission signals. Previous chapters in this thesis have defined the spatial and temporal emission patterns of emissions from natural gas production regions and the use of langrangian models and analysis of ambient concentrations measured at monitoring sites. The present chapter proposes coupling all of these elements into an innovative and effective solution to the attribution and monitoring of methane emissions.

8.4 References

- (1) Konschnik, K., (2014). Shale Gas Development: A Smart Regulation Framework. Environmental Science & Technology.
- (2) Richardson, N., Gottlieb, M., Krupnick., Wiseman, H., (2013). The State of State Shale Gas Regulation. RFF Report, Washington DC.
- (3) U.S. Environmental Protection Agency, Natural Gas STAR Program. Available at <http://epa.gov/gasstar/> (Retrieved 7/3/2014)
- (4) ICF. (2014). Economic Analysis of Methane Emission Reduction Opportunities in the U.S. Onshore Oil and Natural Gas Industries. Prepared for Environmental Defense Fund. Available at http://www.edf.org/sites/default/files/methane_cost_curve_report.pdf (Retrieved 7/3/2014).
- (5) Allen, T. D., Torres, M. V., Sullivan, D., Harrison, M., Hendler, A., Herndon, S. C., et al. (2013). Measurements of Methane Emissions at Natural Gas Production Sites. Proceedings of the National Academy of Sciences.110, 17768-17773.
- (6) Caulton D. R., Shespon P. B., Santoro, R. L., Sparks J. P, Howarth, R. W., Ingraffea, A. R., Cambaliza M. O. L., Sweeney, C., Karion, A., Davis, K. J., Stirm, B. H., Montzla S. A., Miller, B. R. (2014). Toward a better understanding and quantification of methane emissions from shale gas development. Proceedings of the National Academy of Sciences, 111, 6237 – 6242.
- (7) Medlock, K. B. Modeling the implications of expanded US shale gas production. Energy Strategy Reviews 2012, 1, 33-41.
- (8) Rail Road Commission of Texas, Texas Gas Well Gas Production in the Newark, East (Barnett Shale) Field – 1993-2011. Available at http://www.rrc.state.tx.us/barnettshale/NewarkEastField_1993-2011.pdf (accessed 9/11/2012)).

- (9) Nelson, D., Shorter, J., McManus, J., Zahniser, M., (2006) Sub-par-per-billion detection of nitric oxide in air using thermoelectrically cooled mid-infrared quantum cascade laser spectrometer. *Applied Physics B-Lasers and Optics* 75, 343 – 350.
- (10) Jimenez, R., Herndon, D., Shorter, J., Nelson, D., McManus, J., Zahniser, M., (2005). Atmospheric trace gas measurements using dual quantum cascade laser mid-infrared absorption spectrometer. *Novel In-Pane Semiconductor Lasers IV*. Bellingham SPIE-INT Society Optical Engineering. Proc. SPIE 5738, 318 – 331.
- (11) Yacovitch, T., Herndon S., Roscioli, J., Floerchinger C., *et al.*, (2014). Demonstration of an Ethane Spectrometer for Methane Source Identification. *Environmental Science and Technology* 48, 8028-8034.
- (12) Harrison, M.R.; Shires, T.M.; Wessels, J.K.; Cowgill, R. M; Methane Emissions from the Natural Gas Industry, Volumes 1 – 15, Final Report, GRI-94/0257 and EPA-600/R-96-080, Gas Research Institute and US Environmental Protection Agency, June 1996.
- (12) Harrison, M.R.; Shires, T.M.; Wessels, J.K.; Cowgill, R.M.; Methane Emissions from the Natural Gas Industry, EPA/600/SR-96/080 June 1997.
- (13) Kirchgessner, D.A.; Lott, R.A. Cowgill, R.M.; Harrison, M.R.; Shires, T.M. Estimate of Methane Emissions from the U.S. Natural Gas Industry, *Chemosphere*, 35, 1365-1390, 1997.
- (14) McGaughey, G., Sullivan, D., Smith, D., McDonald-Buller, E., Allen, D. (2009) Meteorological Conditions and Associated Temporal Trends in Total Non-Methane Hydrocarbon and Benzene Concentrations in the Corpus Christi Area. The University of Texas at Austin. Available at http://www.utexas.edu/research/ceer/ccaqp/PDF/CCNATReports/Part3_CCNAT_FY2008_2009_Air_Toxics_Conceptual_Model.pdf (Retrieved 7/3/2014)

9 SUMMARY OF CONTRIBUTIONS AND FINDINGS

The appearance of new extraction technologies such as horizontal drilling and hydraulic fracturing have propelled the development of natural gas production, and driven a need for accurate characterizations of air pollutant emissions from natural gas production. Previous emission estimates have been characterized by large uncertainties. This work has addressed emissions of greenhouse gases (GHG, specifically methane), criteria pollutants, and toxic emissions from natural gas production. Both direct and indirect measurements of emissions have been reported, and the spatial and temporal distributions of emissions as well as the role of very high emitting wells and high emitting sources in determining national emissions have been assessed. The major findings from this work are summarized below:

- For virtually all of the sources for which direct measurements were made (e.g. equipment leaks, pneumatic devices, liquids unloading events) a small subset of the sampled devices or wells account for the majority of the emissions.
- For pneumatic devices the subset of high emitters is characterized by devices that behave in a manner inconsistent with the manufacturer's design. Due to the high and relatively continuous emissions, this subset of devices should be readily detectable in leak detection programs; repair or replacement would have a high impact as an emission reduction strategy. Reducing the emissions of the highest emitting devices from their current level to less than 6 scf/h would reduce total emissions from these pneumatic controllers by an average of 160,000 standard cubic feet per year per device.
- For liquids unloading events, emissions depended most strongly on event frequency. Identification of high emitting wells is therefore relatively straight-forward. Emission reduction strategies should focus on the small subset of wells that have a high frequency of unloadings and account for the majority of the emissions from all wells with liquid unloadings. Based on the analyses of different unloading methods (plunger-lift, non-plunger lift), selection of an unloading method for a particular well should take into

consideration the age of the well and potential frequency of unloadings not just in terms of its effectiveness in clearing the well from liquids, but also in the potential reduction of emissions.

- Reconciliation between direct source measurements (bottom-up measurements) and ambient measurements (top-down analyses) was demonstrated by comparing a 20-month time series of hydrocarbon concentration data from a monitoring station in the center of a shale production region (Barnett Shale), with a site by site inventory of emissions for the region. The inventory, adjusted based on recent direct source measurements, was consistent with observations. Hourly and daily variations in the concentrations were explained by variations in meteorology. Evidence of large episodic emissions was not observed in the data set.
- Based on the analyses of hydrocarbon emissions in the Barnett Shale production region, a spatially resolved methane, ethane, and propane emission inventory was developed. This spatially resolved inventory will be used in comparisons with a variety of measurements schemes in the Barnett Shale, including airborne, vehicle, and ground-based measurements.
- Based on all of these analyses, a multi-scale system was designed that would detect and attribute sources of methane emissions in natural gas production regions.

This work has focused on national and regional scale analyses of emissions. Natural gas production from shale gas formations is already in an advanced stage of development in the U.S; with some basins already reaching their peak of production. An area of further research opportunities lies in the application of the array of concepts and tools described in the present work to other countries and regions in the world, where the technological development for the production of natural gas from shale formations is in an early stage.

Collectively, the direct source measurements, and analyses of ambient air pollutant measurements in natural gas production regions reported in this work improve the

estimation, characterization, and methods for monitoring air quality implications of shale gas production.

APPENDICES

A Methane Emissions from Pneumatic Controllers

A.1 Methods for site selection

Goals and overall sampling strategy. Sampling of pneumatic controllers was conducted in four major regions (Appalachian, Gulf Coast, Mid-continent, Rocky Mountain); based on current characterizations of pneumatic controller design types (continuous vent or intermittent vent, on/off or throttling), it was anticipated that several types of controller designs would need to be sampled; it was also anticipated that there would be multiple types of controller service (e.g., separator level control service) that would influence actuation frequency and other parameters that affect emissions. To sample regions, controller design types and controller service types, it was anticipated that approximately 400 devices would need to be sampled.

Selection of Basins Data from EPA's Greenhouse Gas Reporting Program (GHGRP, Reporting Year 2012) were used to identify the Basins where the ten participant companies had reported pneumatic emissions. Based on this distribution of available Basins in which to sample, the study team selected at least two companies to sample in each region (Appalachian, Gulf Coast, Mid-continent, Rocky Mountain), such that all ten participant companies were sampled. Where possible, the study team selected Basins that produced a mix of device design and service types, rather than visiting sites that reported only one device type. The Study Team was solely responsible for the selection of regions and Basins in which to sample.

Once basins and companies to be sampled in each basin were selected, 2-3 day site visits were planned. In each of these site visits, the focus was on sampling well pads for a single company in a single basin. Thirteen of these 2-3 day visits were conducted.

Selection of well pads. Local contacts for participant companies provided descriptions or lists of the well pad sites or central facilities (e.g. sites with separators for multiple wells),

with pneumatic controllers, in the area to be sampled. The study team selected the sites; all sites identified by the local contacts were available for sampling; depending on the distances between sites, the study team either randomly selected sites or selected sites that were relatively proximate to the starting location, so that a relatively large number of samples could be made without losing efficiency due to long travel times between pad locations. The goal was to sample a cross section of typical facilities. If a company had a mix of old and new facilities, or acquired and company built facilities, the study team selected pad types in proportion to the population in the area.

Sampling of devices on well pads or central facilities. Once at a site, the Study Team measured emissions from all pneumatic devices at the site, unless safety or operational issues or lack of access prohibited sampling. This was done to achieve a sample population that would represent controller service and design types currently in use.

A.2 Corrections to Instrument Flow Measurements based on Gas Composition

The supply gas flow meter was calibrated by the manufacturer (Fox Thermal Instruments) using pure methane; instrument flow rates were corrected to account for natural gas compositions, which varied from site to site. Because the flow meter measurement is based on thermal conductivity, the composition correction was based on the relative thermal conductivities of the gas used by pneumatic devices at each site and the pure methane used as a calibration gas.

$$scf/h_{corrected} = scf/h_{inst} \left(\frac{k_{CH_4}}{k_{gas}} \right) \quad (A2.1)$$

where scf/h_{inst} is the raw instrument flow rate reading in standard cubic feet per hour, k_{CH_4} is the pure component thermal conductivity (W/m*K) for methane at standard conditions (70°F and 14.7 psia), and k_{gas} is the thermal conductivity of the gas sampled at standard conditions as determined from the company-provided gas analysis for the site. The

thermal conductivity of the sampled gas (k_{gas}) was calculated as a molar weighted average:

$$k_{gas} = \sum_{i=1}^6 k_i n_i \quad (A2.2)$$

where n_i is the mole fraction of species i in the gas sample for the site and k_i is the pure component thermal conductivity of the species (<http://webbook.nist.gov/chemistry/fluid/>) under standard conditions. For this work, the pure component species considered were methane, ethane, propane, nitrogen, and carbon dioxide. All higher hydrocarbons with a carbon count of four or greater were lumped with butane for purposes of the gas composition correction. The pure component properties used in the calculations are shown in Table A-1. For the range of natural gas compositions from sites visited in this project, the composition correction factor ranged between 1.009 (0.9% increase from raw instrument data) and 1.215 (21.5% increase from raw instrument flow data).

Table A-1. Thermal conductivity (<http://webbook.nist.gov/chemistry/fluid/>) of natural gas components at standard conditions (14.7 psia and 70°F)

Species	Thermal Conductivity (W/m*K)
Methane	0.033759
Ethane	0.020491
Propane	0.017884
Butane +	0.016181
Nitrogen	0.025473
Carbon Dioxide	0.016331

A.3 Field and laboratory comparisons of supply gas and Hi Flow ® measurements

Laboratory measurements

The measurement instruments were laboratory tested using gas flows generated with a mass flow controller. The test flows were designed to mimic the patterns of time varying flow expected from intermittent vent controllers. The flow out of the mass flow

controller was routed through the in-line supply gas meter then fed directly into the feed for the Hi Flow® system (the supply gas meter had a sample frequency of 10 Hz, while the Hi Flow® had a sampling frequency of 0.3 – 0.5 Hz). Tests for the High Flow instrument were completed both with the emissions point bagged and with the emissions being directly fed into the Hi Flow® inlet. In selected experiments, the mass flow controller output was compared to the results of a dry test meter.

Multiple tests were conducted. Figure A-1 shows the results of one test scenario with a pure methane gas stream. The initial flow pattern consisted of three step up-step down flows with a maximum flow of 100 scf/h (a simulated controller actuation). The duration of each step was 30 seconds, and the period between simulated actuations was 2.5 minutes. The second part of the test consisted of three step up-step down flows that also had a maximum flow of 100 scf/h. Each step had a duration of 3 seconds, and the period between simulated actuations was 2.5 minutes.

The time integrated flow per simulated actuation from the mass flow controller was compared to the time integrated flow for the supply gas meter and the Hi Flow® device. The ratio of the integrated supply gas meter flow to the integrated mass flow controller flow averaged 1.07 for this test; the ratio of the integrated Hi Flow® device flow to the integrated mass flow controller flow averaged 0.983.

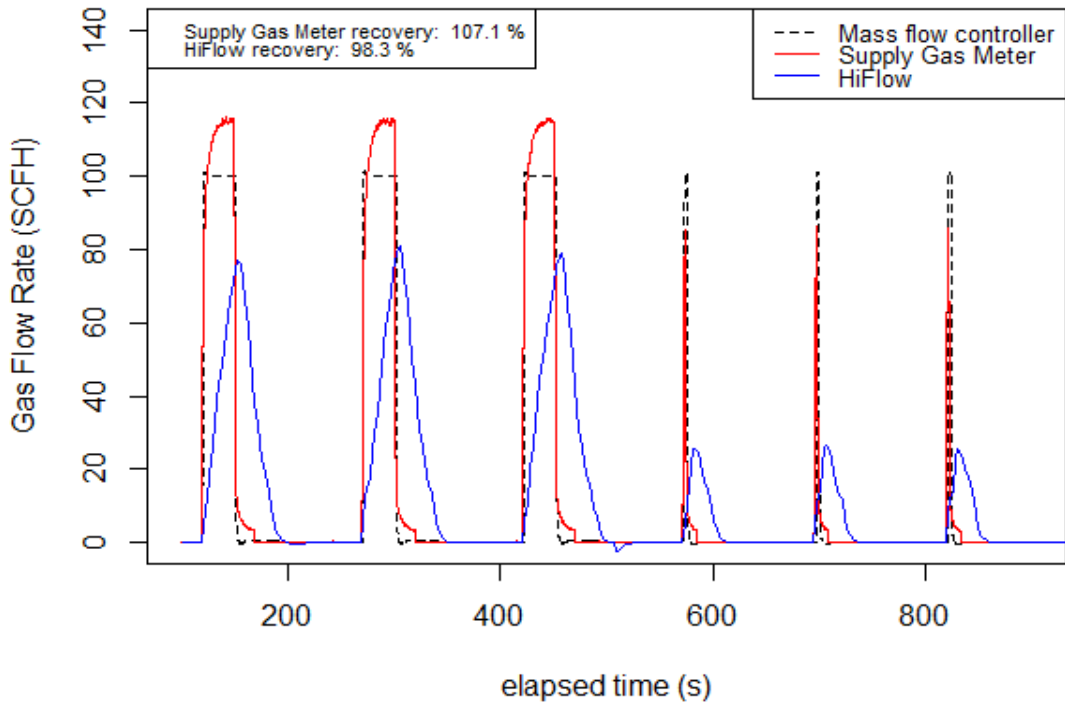


Figure A-1. Laboratory test comparing supply gas meter to Hi Flow® and mass flow controller. The test consisted of two sets of three 100 scf/h simulated actuations; the first set of actuations had a duration of 30 seconds while the second set of actuations had durations of 3 seconds.

For another test (with pure methane test gas), four simulated actuations were performed (30 seconds between actuations), with an increasing maximum flow rate (10, 30, 50, and 100 scf/h, respectively); each simulated actuation had a duration of 20 seconds (Figure A-2). In this test, the ratio of the integrated supply gas meter flow to the integrated mass flow controller flow averaged 1.035; the ratio of the integrated Hi Flow® device flow to the integrated mass flow controller flow averaged 0.952.

Table A-2 summarizes the laboratory tests that were conducted, showing the relative ratio of integrated flows for each case (with pure methane test gas).

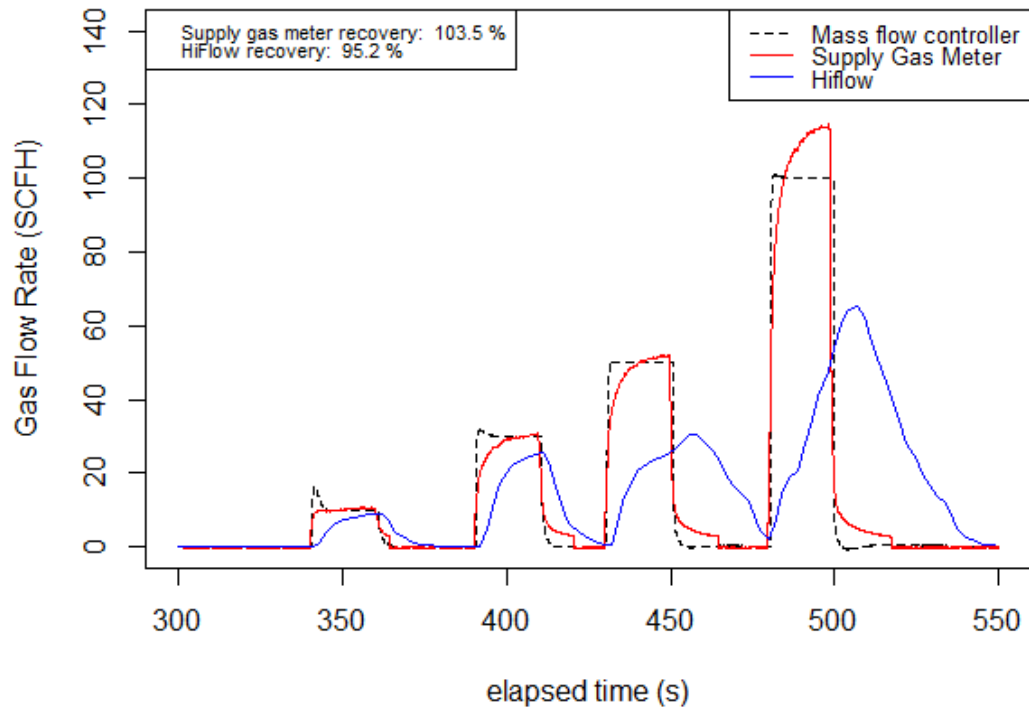


Figure A-2. Laboratory test comparing Supply Gas Meter to Hi Flow® and Mass Flow Controller. The test consisted of four actuations with maximum flows of 10 scf/h, 30 scf/h, 50 scf/h, and 100 scf/h, with a duration of 20 seconds and an interval between actuations of 30 seconds.

Table A-2. Laboratory tests simulating controller actuations, using pure methane, comparing the Supply Gas Meter to the Hi Flow® device.

Test Description	Ratio of integrated supply gas meter flow to integrated mass flow controller flow	Ratio of integrated Hi Flow® device flow to integrated mass flow controller flow
<i>Step up-step down flow. Maximum flow was 100 scf/h, duration of each step was 30 seconds and the period between simulated actuations was 2.5 minutes.</i>	1.07	0.983
<i>Set of actuations with maximum flow of 10 scf/h, 30 scf/h, 50 scf/h, and 100 scf/h with a duration of 20 seconds per actuation, and an interval of 30 seconds between actuations.</i>	1.03	0.952
<i>Set of actuations with a maximum flow of 100 scf/h, 30 scf/h and 10 scf/h, with a duration of 1 second per actuation and an interval of 1 second between actuations.</i>	0.856	0.959
<i>Set of three actuations with a maximum flow of 100 scf/h with a duration of 30 seconds per actuation. Interval of 2.5 minutes between actuations.</i>	1.10	0.922
<i>Set of three actuations with a maximum flow of 100 scf/h with a duration of 3 seconds per actuation. Interval of 2.5 minutes between actuations.</i>	0.829	1.58

Additional tests with a wet gas surrogate (70.5% methane by volume) were completed in the lab before field sampling (Table A-3). For the tests listed in Table A-3, the average ratio of instrument reported mass flow rate to the controller mass flow rate (uncorrected for composition) is 0.878 for the supply gas meter and 0.608 for the Hi Flow®.

Table A-3. Laboratory tests simulating controller actuations, comparing the supply gas meter to the Hi Flow® device with a wet gas surrogate (70.5% methane by volume). Each test consisted of three 30 second bursts at 3 scf/h and at 50 scf/h.

Test Number	Ratio of integrated supply gas meter flow to integrated mass flow controller flow	Ratio of integrated Hi Flow® device flow to integrated mass flow controller flow
<i>1</i>	Not Tested	0.663
<i>2</i>	0.857	0.552
<i>3</i>	0.898	Not Tested

If the supply gas meter is corrected for composition, the average ratio of instrument reported mass flow rate to the controller mass flow rate is 1.034 for the wet gas tests. The Hi Flow® instrument reading is difficult to correct for composition for non-steady flows. At low emission rates, the composition correction would be based on a molar flow rate of carbon, since the sampled gas is catalytically oxidized before it is sent to a thermal conductivity detector. In contrast, at high flow rates, the sample is sent directly to a thermal conductivity detector and the gas composition adjustment would be based on a molar weighted thermal conductivity (see Section A.2). These two adjustments would require precise identification of the transitions between sampling regimes, making the interpretation of field data difficult. Therefore, to adjust Hi Flow® measurements for gas composition in field measurements, the following procedure was used:

1. Determine whether a wet gas or dry gas adjustment will be used; if the percentage methane in the produced gas at a site was greater than 85%, a dry gas correction was used; otherwise a wet gas correction was used
2. For dry gas sites, the Hi Flow® measurement, uncorrected for gas composition, was converted to a supply gas measurement, uncorrected for gas composition. This conversion is based on the laboratory data collected on pure methane reported in Table A-3 (ratio of Supply gas flow rate to Hi Flow® flow rate of 1.12). The uncorrected supply gas flow rate was then corrected for composition using the method described in Section A.2.

3. For wet gas sites, the Hi Flow® measurement, uncorrected for gas composition, was converted to a supply gas measurement, uncorrected for gas composition. This conversion is based on the laboratory data collected on a surrogate wet gas reported in Table A-4 (ratio of Supply gas flow rate to Hi Flow® flow rate of 0.878/0.608; 1.44). The uncorrected supply gas flow rate was then corrected for composition using the method described in Section A.2.

Field Measurements

In field measurements, for 29 controllers, both supply gas and Hi Flow® measurements were made, simultaneously. Four of those measurements will not be analyzed because they occurred at a site (DL02) where the Hi Flow® was capturing a leak signal from equipment, not associated with the controller, located inside the enclosure where the pneumatic devices were installed. An additional measurement (device XQ01-PC04) will not be analyzed because it was concluded that the controller was not completely bagged.

For the final data set of 24 controllers with coupled measurements, 11 (46%) had emissions of less than 0.005 scf/h of whole gas, as measured by the supply gas meter. The results for the remaining 13 devices are shown in Table A-4. For devices with an average emission rate greater than 6 scf/h (measured by the supply gas meter), the supply gas meter to Hi Flow® measurement ratio is between 0.7 and 1.1. These results suggest that for controllers that account for most of the emissions, the two methods produce similar results. For lower flow rates, the lack of agreement is generally due to higher supply gas measurements, relative to Hi Flow® measurements. This is likely due to leaks in the controller/control valve gas line that were not captured by the Hi Flow® enclosure. For the samples where the Hi Flow® device was measuring higher emissions than the supply gas meter, the Hi Flow® may have been sampling emissions that were not in the controller/control valve gas line.

Table A-4. Supply gas meter and Hi Flow® for thirteen controllers where both instruments were used to make measurements.

Device Name	Supply Gas meter - avg. emission rate Whole Gas (scf/h)	Hi Flow® - avg. emission rate Whole Gas (scf/h)	Supply Gas meter to Hi Flow® measurement ratio
AP04-PC02	0.111	0	-
AP04-PC03	0.036	0.007	5.4
AP05-PC01	0.169	0.197	0.9
DL02-PC11	0.528	0.881	0.6
DL02-PC15	1.360	0.696	2.0
DL02-PC42	0.504	0.001	553
LB03-PC01	50.0	55.3	0.9
LB06-PC05	22.5	20.8	1.1
LB07-PC01	36.3	51.3	0.7
LB07-PC04	27.0	33.9	0.8
RB01-PC14	4.16	0.194	21.5
RQ05-PC02	0.008	5.15	0.0
RQ07-PC03	18.2	21.0	0.9

A.4 Pneumatic controller emission data

Tables A-5 to A-9 contain the data collected for each of the 377 measured devices. The first letter of the device code indicates the host company that operated the site where the measurement was performed. The second letter of the device code indicates the basin where the measurement was performed. Letters for these codes were randomly selected, but are used consistently (i.e., specific letters always represent the same company or basin). Figure A-3 shows the boundaries of the geographical regions into which controllers were grouped. Figure A-4 summarizes the application types, well types, and service types sampled population.

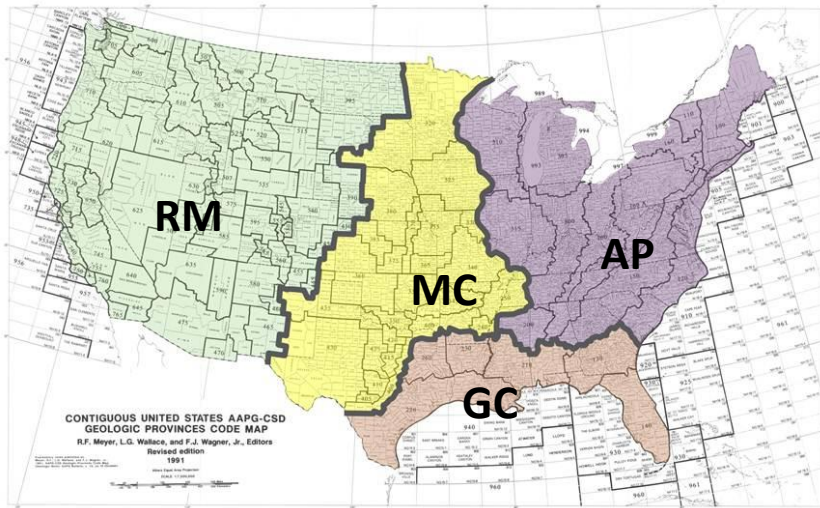


Figure A-3. Measurements of emissions from pneumatic controllers were categorized into Appalachian (AP), Gulf Coast (GC), Mid-Continent (MC) and Rocky Mountain (RM) regions.

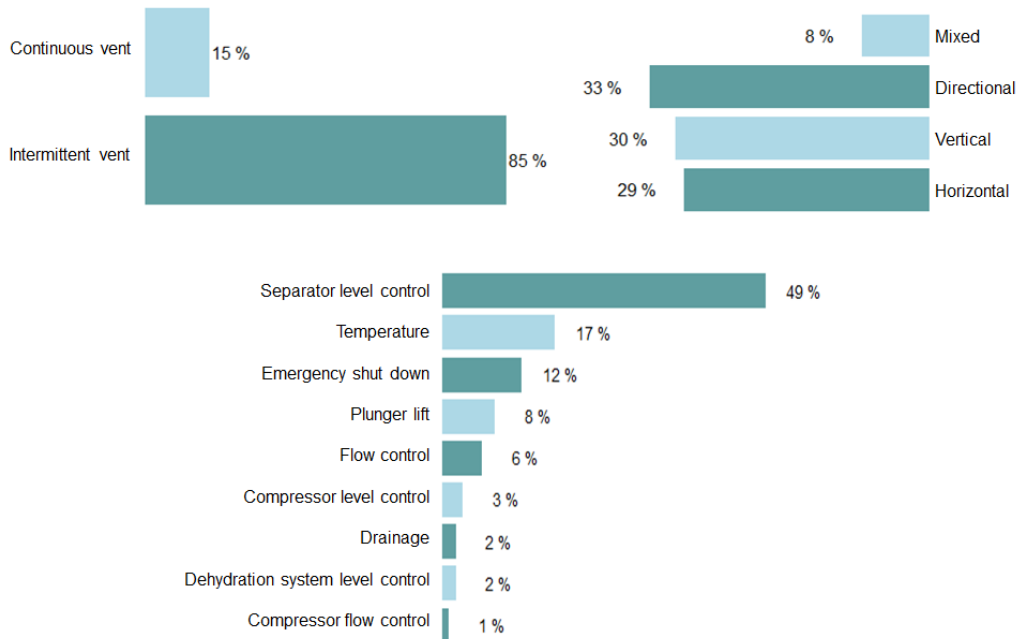


Figure A-4. Sample population characterized by the service type (upper left, continuous vent or intermittent vent), well type (upper right, horizontal, vertical, directional or mixed type), and type of application in which the controller is used (lower, e.g., separator level control).

Table A-5. Descriptive data of measured devices, showing the pneumatic controller application (level, pressure or temperature), process unit they are servicing, and if they are linked to a single well or multiple wells.

Device Identifier	Region	Controller application	Basic application	Detailed application	Single Well (Y/N)
AA01-PC04	MC	Level	Separator	Separator - Level Control	Y
AA01-PC05	MC	Level	Separator	Separator - Level Control	Y
AA01-PC06	MC	Level	Separator	Separator - Level Control	Y
AA01-PC07	MC	Level	Separator	Separator - Level Control	Y
AA01-PC08	MC	Level	Separator	Separator - Level Control	Y
AA01-PC09	MC	Level	Separator	Separator - Level Control	Y
AA01-PC11	MC	Pressure	Compressor	Compressor	Y
AA02-PC04	MC	Level	Compressor	Other	Y
AA02-PC05	MC	Level	Compressor	Compressor	Y
AA02-PC06	MC	Level	Compressor	Compressor	Y
AA02-PC07	MC	Level	Compressor	Compressor	Y
AA02-PC08	MC	Pressure	Compressor	Compressor	Y
AA02-PC09	MC	Level	Compressor	Compressor	Y
AP01-PC01	MC	Level	Separator	Separator - Level Control	N
AP01-PC02	MC	Level	Separator	Separator - Level Control	Y
AP01-PC03	MC	Level	Separator	Separator - Level Control	Y
AP01-PC04	MC	Level	Separator	Separator - Level Control	Y
AP01-PC05	MC	Level	Separator	Separator - Level Control	Y
AP01-PC12	MC	Level	Separator	Separator - Level Control	N
AP02-PC01	MC	Level	Separator	Separator - Level Control	N
AP02-PC02	MC	Level	Separator	Separator - Level Control	N
AP02-PC03	MC	Level	Separator	Separator - Level Control	N
AP02-PC04	MC	Level	Separator	Separator - Level Control	N
AP03-PC01	MC	Level	Separator	Separator - Level Control	Y
AP03-PC02	MC	Level	Separator	Separator - Level Control	N
AP04-PC01	MC	Level	Separator	Separator - Level Control	N
AP04-PC02	MC	Level	Separator	Separator - Level Control	N
AP04-PC03	MC	Level	Separator	Separator - Level Control	N
AP04-PC04	MC	Level	Separator	Separator - Level Control	Y
AP05-PC01	MC	Level	Separator	Separator - Level Control	N
AP05-PC02	MC	Level	Separator	Separator - Level Control	Y
CW01-PC01	RM	Pressure	Wellhead	ESD	Y
CW01-PC02	RM	Pressure	Plunger Lift	Plunger Lift	Y
CW01-PC03	RM	Level	Separator	Separator - Level Control	Y
CW01-PC04	RM	Level	Separator	Separator - Level Control	Y
CW01-PC05	RM	Temperature	Process Heater	Process Heater	Y

Table A-5 (continued)

Device Identifier	Region	Controller application	Basic application	Detailed application	Single Well (Y/N)
CW01-PC11	RM	Pressure	Wellhead	ESD	Y
CW01-PC12	RM	Pressure	Plunger Lift	Plunger Lift	Y
CW01-PC13	RM	Level	Separator	Separator - Level Control	Y
CW01-PC14	RM	Level	Separator	Separator - Level Control	Y
CW01-PC15	RM	Temperature	Process Heater	Process Heater	Y
CW01-PC21	RM	Pressure	Wellhead	ESD	Y
CW01-PC22	RM	Pressure	Plunger Lift	Plunger Lift	Y
CW01-PC23	RM	Level	Separator	Separator - Level Control	Y
CW01-PC24	RM	Level	Separator	Separator - Level Control	Y
CW01-PC25	RM	Temperature	Process Heater	Process Heater	Y
CW01-PC31	RM	Pressure	Wellhead	ESD	Y
CW01-PC32	RM	Pressure	Plunger Lift	Plunger Lift	Y
CW01-PC33	RM	Level	Separator	Separator - Level Control	Y
CW01-PC34	RM	Level	Separator	Separator - Level Control	Y
CW01-PC35	RM	Temperature	Process Heater	Process Heater	Y
CW02-PC01	RM	Pressure	Wellhead	ESD	Y
CW02-PC02	RM	Pressure	Plunger Lift	Plunger Lift	Y
CW02-PC03	RM	Level	Separator	Separator - Level Control	Y
CW02-PC04	RM	Level	Separator	Separator - Level Control	Y
CW02-PC05	RM	Temperature	Process Heater	Process Heater	Y
CW02-PC11	RM	Pressure	Wellhead	ESD	Y
CW02-PC12	RM	Pressure	Plunger Lift	Plunger Lift	Y
CW02-PC13	RM	Level	Separator	Separator - Level Control	Y
CW02-PC14	RM	Level	Separator	Separator - Level Control	Y
CW02-PC15	RM	Temperature	Process Heater	Process Heater	Y
CW02-PC21	RM	Pressure	Wellhead	ESD	Y
CW02-PC22	RM	Pressure	Plunger Lift	Plunger Lift	Y
CW02-PC23	RM	Level	Separator	Separator - Level Control	Y
CW02-PC24	RM	Level	Separator	Separator - Level Control	Y
CW02-PC25	RM	Temperature	Process Heater	Process Heater	Y
CW02-PC31	RM	Pressure	Wellhead	ESD	Y
CW02-PC32	RM	Pressure	Plunger Lift	Plunger Lift	Y
CW02-PC33	RM	Level	Separator	Separator - Level Control	Y
CW02-PC34	RM	Level	Separator	Separator - Level Control	Y
CW02-PC35	RM	Temperature	Process Heater	Process Heater	Y
CZ01-PC01	GC	Level	Separator	Separator - Level Control	Y
CZ01-PC02	GC	Pressure	Plunger Lift	Plunger Lift	Y
CZ02-PC01	GC	Level	Separator	Separator - Level Control	Y

Table A-5 (continued)

Device Identifier	Region	Controller application	Basic application	Detailed application	Single Well (Y/N)
CZ02-PC02	GC	Pressure	Plunger Lift	Plunger Lift	Y
CZ03-PC01	GC	Level	Separator	Separator - Level Control	Y
CZ03-PC02	GC	Pressure	Plunger Lift	Plunger Lift	Y
CZ04-PC01	GC	Level	Separator	Separator - Level Control	Y
CZ04-PC02	GC	Pressure	Plunger Lift	Plunger Lift	Y
CZ05-PC01	GC	Level	Separator	Separator - Level Control	Y
CZ05-PC02	GC	Pressure	Plunger Lift	Plunger Lift	Y
CZ05-PC03	GC	Level	Separator	Separator - Level Control	Y
CZ06-PC01	GC	Level	Separator	Separator - Level Control	Y
CZ06-PC02	GC	Pressure	Plunger Lift	Plunger Lift	Y
CZ06-PC03	GC	Level	Separator	Separator - Level Control	Y
CZ07-PC01	GC	Level	Separator	Separator - Level Control	Y
CZ07-PC02	GC	Pressure	Plunger Lift	Plunger Lift	Y
CZ08-PC01	GC	Level	Separator	Separator - Level Control	Y
CZ08-PC02	GC	Level	Separator	Separator - Level Control	Y
CZ09-PC01	GC	Level	Separator	Separator - Level Control	Y
CZ09-PC02	GC	Level	Separator	Separator - Level Control	Y
CZ10-PC01	GC	Level	Separator	Separator - Level Control	Y
CZ11-PC01	GC	Level	Separator	Separator - Level Control	Y
DL01-PC01	RM	Level	Separator	Separator - Level Control	Y
DL01-PC02	RM	Level	Separator	Separator - Level Control	Y
DL01-PC03	RM	Temperature	Process Heater	Process Heater	Y
DL01-PC04	RM	Level	Wellhead	ESD	Y
DL01-PC05	RM	Temperature	Process Heater	Process Heater	Y
DL01-PC11	RM	Level	Separator	Separator - Level Control	Y
DL01-PC12	RM	Level	Wellhead	ESD	Y
DL01-PC13	RM	Level	Separator	Separator - Level Control	Y
DL01-PC14	RM	Level	Wellhead	ESD	Y
DL01-PC15	RM	Level	Separator	Separator - Level Control	Y
DL01-PC21	RM	Level	Wellhead	ESD	Y
DL01-PC22	RM	Level	Separator	Separator - Level Control	Y
DL01-PC23	RM	Level	Wellhead	ESD	Y
DL01-PC24	RM	Level	Separator	Separator - Level Control	Y
DL01-PC25	RM	Level	Wellhead	ESD	Y
DL01-PC31	RM	Temperature	Process Heater	Process Heater	Y
DL01-PC32	RM	Level	Separator	Separator - Level Control	Y
DL01-PC33	RM	Level	Wellhead	ESD	Y
DL01-PC34	RM	Level	Separator	Separator - Level Control	Y

Table A-5 (continued)

Device Identifier	Region	Controller application	Basic application	Detailed application	Single Well (Y/N)
DL01-PC35	RM	Level	Wellhead	ESD	Y
DL01-PC41	RM	Level	Separator	Separator - Level Control	Y
DL01-PC42	RM	Level	Wellhead	ESD	Y
DL01-PC43	RM	Level	Separator	Separator - Level Control	Y
DL01-PC44	RM	Level	Wellhead	ESD	Y
DL01-PC45	RM	Temperature	Flare	Other	N
DL02-PC01	RM	Level	Separator	Separator - Level Control	Y
DL02-PC02	RM	Level	Wellhead	ESD	Y
DL02-PC03	RM	Level	Separator	Separator - Level Control	Y
DL02-PC04	RM	Level	Wellhead	ESD	Y
DL02-PC05	RM	Level	Separator	Separator - Level Control	Y
DL02-PC11	RM	Level	Wellhead	ESD	Y
DL02-PC12	RM	Level	Separator	Separator - Level Control	Y
DL02-PC13	RM	Level	Wellhead	ESD	Y
DL02-PC14	RM	Level	Separator	Separator - Level Control	Y
DL02-PC15	RM	Level	Wellhead	ESD	Y
DL02-PC21	RM	Level	Separator	Separator - Level Control	Y
DL02-PC22	RM	Level	Wellhead	ESD	Y
DL02-PC23	RM	Level	Separator	Separator - Level Control	Y
DL02-PC24	RM	Level	Wellhead	ESD	Y
DL02-PC25	RM	Level	Separator	Separator - Level Control	Y
DL02-PC31	RM	Level	Wellhead	ESD	Y
DL02-PC32	RM	Level	Separator	Separator - Level Control	Y
DL02-PC33	RM	Level	Wellhead	ESD	Y
DL02-PC34	RM	Level	Separator	Separator - Level Control	Y
DL02-PC35	RM	Level	Wellhead	ESD	Y
DL02-PC41	RM	Level	Separator	Separator - Level Control	Y
DL02-PC42	RM	Level	Wellhead	ESD	Y
DL02-PC43	RM	Level	Separator	Separator - Level Control	Y
DL02-PC44	RM	Level	Wellhead	ESD	Y
DL02-PC45	RM	Level	Separator	Separator - Level Control	Y
DL02-PC51	RM	Level	Wellhead	ESD	Y
DL02-PC52	RM	Level	Separator	Separator - Level Control	Y
DL02-PC53	RM	Level	Wellhead	ESD	Y
DL02-PC54	RM	Temperature	Process Heater	Separator - Other	Y
DL02-PC55	RM	Temperature	Process Heater	Separator - Other	Y
DL02-PC61	RM	Temperature	Process Heater	Separator - Other	Y
DL02-PC62	RM	Temperature	Flare	Other	Y

Table A-5 (continued)

Device Identifier	Region	Controller application	Basic application	Detailed application	Single Well (Y/N)
GZ01-PC01	GC	Level	Separator	Separator - Level Control	Y
GZ01-PC02	GC	Level	Separator	Separator - Level Control	Y
GZ01-PC03	GC	Level	Separator	Separator - Level Control	Y
GZ01-PC04	GC	Level	Separator	Separator - Level Control	Y
GZ01-PC05	GC	Level	Separator	Separator - Level Control	Y
GZ01-PC11	GC	Level	Separator	Separator - Level Control	Y
GZ01-PC12	GC	Level	Separator	Separator - Level Control	Y
GZ01-PC13	GC	Level	Separator	Separator - Level Control	Y
GZ01-PC14	GC	Pressure	Wellhead	ESD	Y
GZ01-PC15	GC	Pressure	Wellhead	ESD	Y
GZ01-PC21	GC	Pressure	Wellhead	ESD	Y
GZ01-PC22	GC	Pressure	Wellhead	ESD	Y
GZ02-PC01	GC	Level	Separator	Separator - Level Control	Y
GZ02-PC02	GC	Level	Separator	Separator - Level Control	Y
GZ02-PC03	GC	Level	Separator	Separator - Level Control	Y
GZ02-PC04	GC	Level	Separator	Separator - Level Control	Y
GZ02-PC05	GC	Level	Compressor	Other	N
GZ02-PC11	GC	Level	Compressor	Compressor	N
GZ02-PC12	GC	Level	Compressor	Compressor	N
GZ02-PC13	GC	Level	Compressor	Compressor	N
GZ02-PC14	GC	Pressure	Compressor	Compressor	N
GZ02-PC15	GC	Level	Compressor	Compressor	N
GZ02-PC21	GC	Level	Compressor	Compressor	N
GZ02-PC22	GC	Pressure	Wellhead	ESD	Y
GZ02-PC23	GC	Pressure	Wellhead	ESD	Y
GZ02-PC24	GC	Pressure	Compressor	Compressor	N
GZ03-PC01	GC	Level	Compressor	Compressor	N
GZ03-PC02	GC	Level	Compressor	Compressor	N
GZ03-PC03	GC	Level	Compressor	Compressor	N
GZ03-PC04	GC	Level	Compressor	Other	N
GZ03-PC05	GC	Pressure	Compressor	Compressor	N
GZ03-PC11	GC	Pressure	Wellhead	ESD	Y
GZ03-PC12	GC	Pressure	Wellhead	ESD	Y
GZ03-PC13	GC	Level	Separator	Separator - Level Control	N
GZ03-PC14	GC	Level	Separator	Separator - Level Control	N
GZ03-PC15	GC	Level	Separator	Separator - Level Control	N
GZ03-PC21	GC	Level	Separator	Separator - Level Control	Y
GZ03-PC22	GC	Level	Separator	Separator - Level Control	Y

Table A-5 (continued).

Device Identifier	Region	Controller application	Basic application	Detailed application	Single Well (Y/N)
GZ03-PC23	GC	Level	Separator	Separator - Level Control	Y
GZ03-PC24	GC	Level	Separator	Separator - Level Control	Y
GZ03-PC25	GC	Pressure	Wellhead	ESD	N
GZ03-PC31	GC	Level	Dehydration System	Separator - Level Control	N
GZ03-PC32	GC	Level	Dehydration System	Dehydration System	N
GZ03-PC33	GC	Level	Dehydration System	Dehydration System	N
GZ03-PC34	GC	Temperature	Dehydration System	Dehydration System	N
GZ03-PC35	GC	Temperature	Dehydration System	Dehydration System	N
GZ03-PC41	GC	Level	Dehydration System	Other	N
GZ03-PC42	GC	Pressure	Sales	Other	N
GZ04-PC01	GC	Level	Separator	Separator - Level Control	Y
GZ04-PC02	GC	Level	Separator	Separator - Level Control	Y
GZ04-PC03	GC	Level	Separator	Separator - Level Control	Y
GZ04-PC04	GC	Level	Separator	Separator - Level Control	Y
GZ04-PC05	GC	Pressure	Wellhead	ESD	Y
GZ04-PC11	GC	Pressure	Wellhead	ESD	Y
LB01-PC01	GC	Level	Separator	Separator - Level Control	Y
LB01-PC02	GC	Level	Separator	Separator - Level Control	Y
LB02-PC01	GC	Level	Separator	Separator - Level Control	Y
LB02-PC02	GC	Level	Separator	Separator - Level Control	Y
LB02-PC03	GC	Level	Separator	Separator - Level Control	Y
LB02-PC04	GC	Level	Separator	Separator - Level Control	Y
LB03-PC01	GC	Level	Separator	Separator - Level Control	Y
LB03-PC02	GC	Level	Separator	Separator - Level Control	Y
LB04-PC01	GC	Level	Separator	Separator - Level Control	Y
LB04-PC02	GC	Level	Separator	Separator - Level Control	Y
LB04-PC03	GC	Level	Separator	Separator - Level Control	Y
LB04-PC04	GC	Level	Separator	Separator - Level Control	Y
LB05-PC01	GC	Level	Separator	Separator - Level Control	Y
LB05-PC02	GC	Level	Separator	Separator - Level Control	Y
LB05-PC03	GC	Level	Separator	Separator - Level Control	Y
LB05-PC04	GC	Level	Separator	Separator - Level Control	Y
LB06-PC01	GC	Level	Separator	Separator - Level Control	Y
LB06-PC02	GC	Level	Separator	Separator - Level Control	Y
LB06-PC03	GC	Level	Separator	Separator - Level Control	Y
LB06-PC04	GC	Level	Separator	Separator - Level Control	Y
LB06-PC05	GC	Level	Separator	Separator - Level Control	Y
LB06-PC11	GC	Level	Separator	Separator - Level Control	Y

Table A-5 (continued).

Device Identifier	Region	Controller application	Basic application	Detailed application	Single Well (Y/N)
LB07-PC01	GC	Level	Separator	Separator - Level Control	Y
LB07-PC02	GC	Level	Separator	Separator - Level Control	Y
LB07-PC03	GC	Level	Separator	Separator - Level Control	Y
LB07-PC04	GC	Level	Separator	Separator - Level Control	Y
OF01-PC01	AP	Pressure	Wellhead	Separator - Other	Y
OF02-PC01	AP	Pressure	Wellhead	Separator - Other	Y
OF03-PC01	AP	Pressure	Wellhead	Separator - Other	Y
OF04-PC01	AP	Pressure	Wellhead	Separator - Other	Y
OF05-PC01	AP	Pressure	Wellhead	Separator - Other	Y
OF06-PC01	AP	Pressure	Wellhead	Separator - Other	Y
OF07-PC01	AP	Pressure	Wellhead	Separator - Other	Y
OF08-PC01	AP	Pressure	Wellhead	Separator - Other	Y
OF08-PC02	AP	Pressure	Plunger Lift	Plunger Lift	Y
OF09-PC01	AP	Pressure	Wellhead	Separator - Other	Y
OF10-PC01	AP	Pressure	Wellhead	Separator - Other	Y
OF11-PC01	AP	Pressure	Wellhead	Separator - Other	Y
OF11-PC02	AP	Level	Separator	Separator - Level Control	Y
OF11-PC03	AP	Level	Separator	Separator - Level Control	Y
OF11-PC04	AP	Pressure	Wellhead	Separator - Other	Y
RB01-PC01	GC	Level	Separator	Separator - Level Control	Y
RB01-PC02	GC	Level	Separator	Separator - Level Control	Y
RB01-PC03	GC	Level	Separator	Separator - Level Control	Y
RB01-PC04	GC	Pressure	Separator	Separator - Other	N
RB01-PC05	GC	Level	Separator	Separator - Level Control	N
RB01-PC11	GC	Level	Separator	Separator - Level Control	N
RB01-PC12	GC	Level	Separator	Separator - Level Control	N
RB01-PC13	GC	Level	Dehydration System	Dehydration System	N
RB01-PC14	MC	Level	Dehydration System	Dehydration System	N
RB01-PC15	GC	Level	Dehydration System	Dehydration System	N
RB01-PC21	GC	Temperature	Dehydration System	Dehydration System	N
RB01-PC23	GC	Pressure	Dehydration System	Other	N
RB01-PC24	GC	Level	Dehydration System	Separator - Level Control	N
RB01-PC25	GC	Level	Flare	Other	N
RB01-PC34	GC	Level	Separator	Separator - Level Control	Y
RB02-PC01	GC	Level	Separator	Separator - Level Control	Y
RB02-PC02	GC	Level	Dehydration System	Dehydration System	Y
RB02-PC03	GC	Temperature	Dehydration System	Dehydration System	Y
RB02-PC05	GC	Level	Dehydration System	Dehydration System	Y

Table A-5 (continued).

Device Identifier	Region	Controller application	Basic application	Detailed application	Single Well (Y/N)
RB02-PC11	GC	Level	Dehydration System	Dehydration System	Y
RB02-PC12	GC	Level	Dehydration System	Other	Y
RQ01-PC01	MC	Level	Separator	Separator - Level Control	Y
RQ02-PC01	MC	Level	Separator	Separator - Level Control	Y
RQ02-PC02	MC	Pressure	Plunger Lift	Plunger Lift	Y
RQ03-PC01	MC	Level	Separator	Separator - Level Control	Y
RQ04-PC01	MC	Level	Separator	Separator - Level Control	Y
RQ05-PC01	MC	Level	Separator	Separator - Level Control	Y
RQ05-PC02	MC	Pressure	Plunger Lift	Plunger Lift	Y
RQ05-PC03	MC	Level	Separator	Separator - Level Control	Y
RQ05-PC04	MC	Pressure	Plunger Lift	Plunger Lift	Y
RQ06-PC01	MC	Temperature	Process Heater	Process Heater	Y
RQ06-PC02	MC	Temperature	Process Heater	Process Heater	Y
RQ06-PC03	MC	Level	Separator	Separator - Level Control	Y
RQ06-PC04	MC	Level	Separator	Separator - Level Control	Y
RQ07-PC01	MC	Temperature	Process Heater	Process Heater	Y
RQ07-PC02	MC	Temperature	Process Heater	Process Heater	Y
RQ07-PC03	MC	Level	Separator	Separator - Level Control	Y
RQ07-PC04	MC	Level	Separator	Separator - Level Control	Y
RQ07-PC05	MC	Pressure	Plunger Lift	Plunger Lift	Y
VF01-PC01	AP	Level	Wellhead	ESD	Y
VF01-PC02	AP	Level	Separator	Separator - Level Control	Y
VF01-PC03	AP	Temperature	Process Heater	Process Heater	Y
VF01-PC04	AP	Temperature	Process Heater	Process Heater	Y
VF01-PC05	AP	Level	Wellhead	ESD	Y
VF01-PC11	AP	Level	Separator	Separator - Level Control	Y
VF01-PC12	AP	Temperature	Process Heater	Process Heater	Y
VF01-PC13	AP	Temperature	Process Heater	Process Heater	Y
VF01-PC14	AP	Level	Wellhead	ESD	Y
VF01-PC15	AP	Level	Separator	Separator - Level Control	Y
VF01-PC21	AP	Temperature	Process Heater	Process Heater	Y
VF01-PC22	AP	Temperature	Process Heater	Process Heater	Y
VF01-PC23	AP	Level	Wellhead	ESD	Y
VF01-PC24	AP	Level	Separator	Separator - Level Control	Y
VF01-PC25	AP	Temperature	Process Heater	Process Heater	Y
VF01-PC31	AP	Temperature	Process Heater	Process Heater	Y
VF01-PC32	AP	Level	Wellhead	ESD	Y
VF01-PC33	AP	Level	Separator	Separator - Level Control	Y

Table A-5 (continued).

Device Identifier	Region	Controller application	Basic application	Detailed application	Single Well (Y/N)
VF01-PC34	AP	Temperature	Process Heater	Process Heater	Y
VF01-PC35	AP	Temperature	Process Heater	Process Heater	Y
VF01-PC41	AP	Level	Wellhead	ESD	Y
VF01-PC42	AP	Level	Separator	Separator - Level Control	Y
VF01-PC43	AP	Temperature	Process Heater	Process Heater	Y
VF01-PC44	AP	Temperature	Process Heater	Process Heater	Y
VF01-PC45	AP	Pressure	Wellhead	Other	Y
VF01-PC51	AP	Pressure	Wellhead	Other	Y
VF01-PC52	AP	Pressure	Wellhead	Other	Y
VF01-PC53	AP	Pressure	Wellhead	Other	Y
VF01-PC54	AP	Pressure	Wellhead	Other	Y
VF01-PC55	AP	Pressure	Wellhead	Other	Y
VF02-PC01	AP	Level	Separator	Separator - Level Control	Y
VF02-PC02	AP	Level	Separator	Separator - Level Control	Y
VF02-PC03	AP	Level	Separator	Separator - Level Control	Y
VF02-PC04	AP	Level	Separator	Separator - Level Control	Y
VF02-PC05	AP	Level	Separator	Separator - Level Control	Y
VF02-PC11	AP	Level	Separator	Separator - Level Control	Y
VF02-PC12	AP	Temperature	Process Heater	Process Heater	Y
XQ01-PC01	MC	Level	Separator	Separator - Level Control	Y
XQ01-PC02	MC	Level	Separator	Separator - Level Control	Y
XQ01-PC03	MC	Temperature	Process Heater	Process Heater	Y
XQ01-PC04	MC	Pressure	Plunger Lift	Plunger Lift	Y
XQ02-PC01	MC	Level	Separator	Separator - Level Control	Y
XQ02-PC02	MC	Level	Separator	Separator - Level Control	Y
XQ02-PC03	MC	Temperature	Process Heater	Process Heater	Y
XQ02-PC04	MC	Pressure	Plunger Lift	Plunger Lift	Y
XQ03-PC01	MC	Level	Separator	Separator - Level Control	Y
XQ03-PC02	MC	Level	Separator	Separator - Level Control	Y
XQ03-PC03	MC	Temperature	Process Heater	Process Heater	Y
XQ03-PC04	MC	Pressure	Plunger Lift	Plunger Lift	Y
XQ04-PC01	MC	Level	Separator	Separator - Level Control	Y
XQ04-PC02	MC	Level	Separator	Separator - Level Control	Y
XQ04-PC03	MC	Temperature	Process Heater	Process Heater	Y
XQ04-PC04	MC	Pressure	Plunger Lift	Plunger Lift	Y
XQ05-PC01	MC	Level	Separator	Separator - Level Control	Y
XQ05-PC02	MC	Level	Separator	Separator - Level Control	Y
XQ05-PC03	MC	Temperature	Process Heater	Process Heater	Y

Table A-5 (continued).

Device Identifier	Region	Controller application	Basic application	Detailed application	Single Well (Y/N)
XQ05-PC04	MC	Pressure	Plunger Lift	Plunger Lift	Y
XQ06-PC01	MC	Level	Separator	Separator - Level Control	Y
XQ06-PC02	MC	Level	Separator	Separator - Level Control	Y
XQ06-PC03	MC	Temperature	Process Heater	Process Heater	Y
XQ06-PC04	MC	Pressure	Plunger Lift	Plunger Lift	Y
XQ07-PC01	MC	Level	Separator	Separator - Level Control	Y
XQ07-PC02	MC	Level	Separator	Separator - Level Control	Y
XQ07-PC03	MC	Temperature	Process Heater	Process Heater	Y
XQ07-PC04	MC	Pressure	Plunger Lift	Plunger Lift	Y
ZW01-PC01	RM	Level	Separator	Separator - Level Control	Y
ZW01-PC02	RM	Level	Separator	Separator - Level Control	Y
ZW01-PC03	RM	Temperature	Dehydration System	Dehydration System	Y
ZW01-PC04	RM	Temperature	Dehydration System	Dehydration System	Y
ZW01-PC05	RM	Temperature	Process Heater	Process Heater	Y
ZW01-PC11	RM	Temperature	Process Heater	Process Heater	Y
ZW01-PC12	RM	Level	Separator	Separator - Level Control	Y
ZW01-PC13	RM	Level	Separator	Separator - Level Control	Y
ZW01-PC14	RM	Level	Separator	Separator - Level Control	Y
ZW01-PC15	RM	Level	Separator	Separator - Level Control	Y
ZW01-PC21	RM	Level	Process Heater	Process Heater	Y
ZW01-PC22	RM	Temperature	Dehydration System	Dehydration System	Y
ZW01-PC23	RM	Temperature	Dehydration System	Dehydration System	Y
ZW01-PC24	RM	Temperature	Process Heater	Process Heater	Y
ZW01-PC25	RM	Temperature	Process Heater	Process Heater	Y
ZW01-PC31	RM	Pressure	Plunger Lift	Plunger Lift	Y
ZW01-PC32	RM	Temperature	Dehydration System	Dehydration System	Y
ZW01-PC33	RM	Temperature	Dehydration System	Dehydration System	Y
ZW01-PC34	RM	Temperature	Process Heater	Process Heater	Y
ZW01-PC35	RM	Temperature	Process Heater	Process Heater	Y
ZW02-PC01	RM	Level	Separator	Separator - Level Control	Y
ZW02-PC02	RM	Level	Separator	Separator - Level Control	Y
ZW02-PC03	RM	Temperature	Dehydration System	Dehydration System	Y
ZW02-PC04	RM	Temperature	Dehydration System	Dehydration System	Y
ZW02-PC12	RM	Pressure	Plunger Lift	Plunger Lift	Y
ZW03-PC01	RM	Level	Separator	Separator - Level Control	Y
ZW03-PC02	RM	Level	Separator	Separator - Level Control	Y
ZW03-PC03	RM	Pressure	Plunger Lift	Plunger Lift	Y

Table A-6. Device classification, based on field characterization by site operator (On/OFF or throttle), time series characterization (Intermittent-vent, Continuous bleed, Intermittent-vent*, or Continuous bleed*, and EPA classification provided by the companies.

Device Identifier	Field Characterization of Device service	Classification based on time series ¹	Company classification into EPA categories ²
AA01-PC04	On/Off	intermittent-vent	intermittent
AA01-PC05	Throttle	intermittent-vent	intermittent
AA01-PC06	On/Off	intermittent-vent	intermittent
AA01-PC07	Throttle	intermittent-vent	intermittent
AA01-PC08	On/Off	intermittent-vent	intermittent
AA01-PC09	Throttle	intermittent-vent	intermittent
AA01-PC11	Throttle	intermittent-vent	intermittent
AA02-PC04	On/Off	continuous bleed	intermittent
AA02-PC05	On/Off	intermittent-vent	intermittent
AA02-PC06	On/Off	continuous bleed *	intermittent
AA02-PC07	On/Off	continuous bleed *	intermittent
AA02-PC08	Throttle	continuous bleed	intermittent
AA02-PC09	On/Off	continuous bleed	intermittent
AP01-PC01	On/Off	intermittent-vent	intermittent
AP01-PC02	On/Off	intermittent-vent	intermittent
AP01-PC03	On/Off	intermittent-vent	intermittent
AP01-PC04	On/Off	intermittent-vent	intermittent
AP01-PC05	On/Off	intermittent-vent	intermittent
AP01-PC12	On/Off	intermittent-vent *	intermittent
AP02-PC01	On/Off	intermittent-vent	intermittent
AP02-PC02	On/Off	intermittent-vent	intermittent
AP02-PC03	On/Off	intermittent-vent	intermittent
AP02-PC04	On/Off	intermittent-vent	intermittent
AP03-PC01	On/Off	intermittent-vent	intermittent
AP03-PC02	On/Off	intermittent-vent	intermittent
AP04-PC01	On/Off	intermittent-vent	intermittent
AP04-PC02	On/Off	intermittent-vent	intermittent
AP04-PC03	On/Off	intermittent-vent	intermittent
AP04-PC04	On/Off	intermittent-vent	intermittent
AP05-PC01	On/Off	intermittent-vent	intermittent
AP05-PC02	On/Off	intermittent-vent	intermittent
CW01-PC01	On/Off	intermittent-vent	low bleed
CW01-PC02	On/Off	intermittent-vent	low bleed
CW01-PC03	On/Off	intermittent-vent	low bleed

Table A-6 (continued).

Device Identifier	Field Characterization of Device service	Classification based on time series ¹	Company classification into EPA categories ²
CW01-PC04	On/Off	intermittent-vent	low bleed
CW01-PC05	On/Off	intermittent-vent	low bleed
CW01-PC11	On/Off	intermittent-vent	low bleed
CW01-PC12	On/Off	intermittent-vent	low bleed
CW01-PC13	On/Off	intermittent-vent	low bleed
CW01-PC14	On/Off	intermittent-vent	low bleed
CW01-PC15	On/Off	intermittent-vent	low bleed
CW01-PC21	On/Off	intermittent-vent	low bleed
CW01-PC22	On/Off	intermittent-vent	low bleed
CW01-PC23	On/Off	intermittent-vent	low bleed
CW01-PC24	On/Off	intermittent-vent	low bleed
CW01-PC25	On/Off	intermittent-vent	low bleed
CW01-PC31	On/Off	intermittent-vent	low bleed
CW01-PC32	On/Off	intermittent-vent	low bleed
CW01-PC33	On/Off	intermittent-vent	low bleed
CW01-PC34	On/Off	intermittent-vent	low bleed
CW01-PC35	On/Off	intermittent-vent	low bleed
CW02-PC01	On/Off	intermittent-vent	intermittent
CW02-PC02	On/Off	intermittent-vent	intermittent
CW02-PC03	On/Off	intermittent-vent	intermittent
CW02-PC04	On/Off	intermittent-vent	intermittent
CW02-PC05	On/Off	intermittent-vent	intermittent
CW02-PC11	On/Off	intermittent-vent	intermittent
CW02-PC12	On/Off	intermittent-vent	intermittent
CW02-PC13	On/Off	intermittent-vent	intermittent
CW02-PC14	On/Off	intermittent-vent	intermittent
CW02-PC15	On/Off	intermittent-vent	intermittent
CW02-PC21	On/Off	intermittent-vent	intermittent
CW02-PC22	On/Off	intermittent-vent	intermittent
CW02-PC23	On/Off	intermittent-vent	intermittent
CW02-PC24	On/Off	intermittent-vent	intermittent
CW02-PC25	On/Off	intermittent-vent	intermittent
CW02-PC31	On/Off	intermittent-vent	intermittent
CW02-PC32	On/Off	intermittent-vent	intermittent
CW02-PC33	On/Off	continuous bleed *	intermittent
CW02-PC34	On/Off	intermittent-vent	intermittent
CW02-PC35	On/Off	intermittent-vent	intermittent

Table A-6 (continued).

Device Identifier	Field Characterization of Device service	Classification based on time series ¹	Company classification into EPA categories ²
CZ01-PC01	On/Off	intermittent-vent	not classified
CZ01-PC02	On/Off	intermittent-vent	not classified
CZ02-PC01	On/Off	continuous bleed *	not classified
CZ02-PC02	On/Off	intermittent-vent	not classified
CZ03-PC01	Throttle	intermittent-vent	not classified
CZ03-PC02	On/Off	intermittent-vent	not classified
CZ04-PC01	On/Off	intermittent-vent	not classified
CZ04-PC02	On/Off	intermittent-vent	not classified
CZ05-PC01	On/Off	continuous bleed	not classified
CZ05-PC02	On/Off	intermittent-vent	not classified
CZ05-PC03	On/Off	intermittent-vent	not classified
CZ06-PC01	Throttle	continuous bleed	not classified
CZ06-PC02	On/Off	continuous bleed	not classified
CZ06-PC03	Throttle	intermittent-vent	not classified
CZ07-PC01	On/Off	intermittent-vent	not classified
CZ07-PC02	On/Off	intermittent-vent	not classified
CZ08-PC01	Throttle	continuous bleed	low bleed
CZ08-PC02	Throttle	intermittent-vent *	low bleed
CZ09-PC01	On/Off	continuous bleed	low bleed
CZ09-PC02	Throttle	continuous bleed	low bleed
CZ10-PC01	On/Off	intermittent-vent *	low bleed
CZ11-PC01	Throttle	intermittent-vent *	low bleed
DL01-PC01	On/Off	intermittent-vent	not classified
DL01-PC02	On/Off	intermittent-vent	not classified
DL01-PC03	On/Off	intermittent-vent	not classified
DL01-PC04	On/Off	continuous bleed *	not classified
DL01-PC05	On/Off	intermittent-vent	not classified
DL01-PC11	On/Off	intermittent-vent	not classified
DL01-PC12	On/Off	intermittent-vent	not classified
DL01-PC13	On/Off	intermittent-vent	not classified
DL01-PC14	On/Off	intermittent-vent	not classified
DL01-PC15	On/Off	intermittent-vent	not classified
DL01-PC21	On/Off	intermittent-vent	not classified
DL01-PC22	On/Off	intermittent-vent	not classified
DL01-PC23	On/Off	intermittent-vent *	not classified
DL01-PC24	On/Off	continuous bleed	not classified
DL01-PC25	On/Off	intermittent-vent	not classified

Table A-6 (continued).

Device Identifier	Field Characterization of Device service	Classification based on time series ¹	Company classification into EPA categories ²
DL01-PC31	On/Off	intermittent-vent	not classified
DL01-PC32	On/Off	intermittent-vent	not classified
DL01-PC33	On/Off	intermittent-vent	not classified
DL01-PC34	On/Off	intermittent-vent	not classified
DL01-PC35	On/Off	intermittent-vent	not classified
DL01-PC41	On/Off	intermittent-vent	not classified
DL01-PC42	On/Off	intermittent-vent	not classified
DL01-PC43	On/Off	intermittent-vent	not classified
DL01-PC44	On/Off	intermittent-vent	not classified
DL01-PC45	On/Off	intermittent-vent	not classified
DL02-PC01	On/Off	intermittent-vent	not classified
DL02-PC02	On/Off	intermittent-vent	not classified
DL02-PC03	On/Off	intermittent-vent	not classified
DL02-PC04	On/Off	intermittent-vent	not classified
DL02-PC05	On/Off	intermittent-vent	not classified
DL02-PC11	On/Off	intermittent-vent	not classified
DL02-PC12	On/Off	intermittent-vent	not classified
DL02-PC13	On/Off	intermittent-vent	not classified
DL02-PC14	On/Off	intermittent-vent *	not classified
DL02-PC15	On/Off	intermittent-vent *	not classified
DL02-PC21	On/Off	intermittent-vent	not classified
DL02-PC22	On/Off	intermittent-vent	not classified
DL02-PC23	On/Off	intermittent-vent	not classified
DL02-PC24	On/Off	intermittent-vent	not classified
DL02-PC25	On/Off	continuous bleed *	not classified
DL02-PC31	On/Off	intermittent-vent	not classified
DL02-PC32	On/Off	intermittent-vent	not classified
DL02-PC33	On/Off	intermittent-vent *	not classified
DL02-PC34	On/Off	intermittent-vent	not classified
DL02-PC35	On/Off	intermittent-vent	not classified
DL02-PC41	On/Off	intermittent-vent	not classified
DL02-PC42	On/Off	intermittent-vent	not classified
DL02-PC43	On/Off	intermittent-vent	not classified
DL02-PC44	On/Off	intermittent-vent	not classified
DL02-PC45	On/Off	intermittent-vent	not classified
DL02-PC51	On/Off	intermittent-vent	not classified
DL02-PC52	On/Off	intermittent-vent	not classified

Table A-6 (continued).

Device Identifier	Field Characterization of Device service	Classification based on time series ¹	Company classification into EPA categories ²
DL02-PC53	On/Off	intermittent-vent	not classified
DL02-PC54	On/Off	intermittent-vent	not classified
DL02-PC55	On/Off	intermittent-vent	not classified
DL02-PC61	On/Off	intermittent-vent	not classified
DL02-PC62	On/Off	intermittent-vent	not classified
GZ01-PC01	On/Off	intermittent-vent	intermittent
GZ01-PC02	On/Off	intermittent-vent	intermittent
GZ01-PC03	On/Off	intermittent-vent	intermittent
GZ01-PC04	On/Off	intermittent-vent	intermittent
GZ01-PC05	On/Off	intermittent-vent	intermittent
GZ01-PC11	On/Off	intermittent-vent	intermittent
GZ01-PC12	On/Off	intermittent-vent	intermittent
GZ01-PC13	On/Off	intermittent-vent	intermittent
GZ01-PC14	On/Off	intermittent-vent	intermittent
GZ01-PC15	On/Off	intermittent-vent	intermittent
GZ01-PC21	On/Off	intermittent-vent	intermittent
GZ01-PC22	On/Off	intermittent-vent	intermittent
GZ02-PC01	Throttle	intermittent-vent	intermittent
GZ02-PC02	Throttle	intermittent-vent	intermittent
GZ02-PC03	Throttle	intermittent-vent	intermittent
GZ02-PC04	Throttle	intermittent-vent	intermittent
GZ02-PC05	On/Off	intermittent-vent	intermittent
GZ02-PC11	On/Off	intermittent-vent	intermittent
GZ02-PC12	On/Off	continuous bleed	intermittent
GZ02-PC13	On/Off	intermittent-vent	intermittent
GZ02-PC14	Throttle	continuous bleed	intermittent
GZ02-PC15	On/Off	intermittent-vent	intermittent
GZ02-PC21	Throttle	intermittent-vent	intermittent
GZ02-PC22	On/Off	intermittent-vent	intermittent
GZ02-PC23	On/Off	intermittent-vent	intermittent
GZ02-PC24	On/Off	intermittent-vent	intermittent
GZ03-PC01	On/Off	continuous bleed *	intermittent
GZ03-PC02	On/Off	intermittent-vent *	intermittent
GZ03-PC03	On/Off	intermittent-vent	intermittent
GZ03-PC04	On/Off	intermittent-vent	intermittent
GZ03-PC05	Throttle	intermittent-vent *	intermittent
GZ03-PC11	On/Off	intermittent-vent	intermittent

Table A-6 (continued).

Device Identifier	Field Characterization of Device service	Classification based on time series ¹	Company classification into EPA categories ²
GZ03-PC12	On/Off	intermittent-vent	intermittent
GZ03-PC13	Throttle	continuous bleed	intermittent
GZ03-PC14	On/Off	intermittent-vent *	intermittent
GZ03-PC15	On/Off	intermittent-vent	intermittent
GZ03-PC21	On/Off	intermittent-vent	intermittent
GZ03-PC22	On/Off	continuous bleed	intermittent
GZ03-PC23	On/Off	intermittent-vent	intermittent
GZ03-PC24	On/Off	continuous bleed	intermittent
GZ03-PC25	On/Off	intermittent-vent	intermittent
GZ03-PC31	On/Off	intermittent-vent	intermittent
GZ03-PC32	Throttle	intermittent-vent	intermittent
GZ03-PC33	Throttle	intermittent-vent	intermittent
GZ03-PC34	On/Off	intermittent-vent	intermittent
GZ03-PC35	On/Off	intermittent-vent	intermittent
GZ03-PC41	On/Off	intermittent-vent	intermittent
GZ03-PC42	Throttle	intermittent-vent	intermittent
GZ04-PC01	On/Off	intermittent-vent	intermittent
GZ04-PC02	Throttle	continuous bleed *	intermittent
GZ04-PC03	On/Off	intermittent-vent *	intermittent
GZ04-PC04	Throttle	intermittent-vent	intermittent
GZ04-PC05	On/Off	intermittent-vent	intermittent
GZ04-PC11	On/Off	intermittent-vent	intermittent
LB01-PC01	On/Off	intermittent-vent	not classified
LB01-PC02	On/Off	intermittent-vent	not classified
LB02-PC01	On/Off	intermittent-vent	not classified
LB02-PC02	On/Off	intermittent-vent	not classified
LB02-PC03	On/Off	intermittent-vent	not classified
LB02-PC04	On/Off	intermittent-vent	not classified
LB03-PC01	On/Off	intermittent-vent *	not classified
LB03-PC02	On/Off	intermittent-vent	not classified
LB04-PC01	On/Off	intermittent-vent	not classified
LB04-PC02	On/Off	intermittent-vent *	not classified
LB04-PC03	On/Off	intermittent-vent	not classified
LB04-PC04	On/Off	intermittent-vent	not classified
LB05-PC01	On/Off	intermittent-vent *	not classified
LB05-PC02	On/Off	intermittent-vent	not classified
LB05-PC03	On/Off	intermittent-vent *	not classified

Table A-6 (continued).

Device Identifier	Field Characterization of Device service	Classification based on time series ¹	Company classification into EPA categories ²
LB05-PC04	On/Off	intermittent-vent	not classified
LB06-PC01	On/Off	intermittent-vent	not classified
LB06-PC02	On/Off	intermittent-vent	not classified
LB06-PC03	On/Off	intermittent-vent	not classified
LB06-PC04	On/Off	intermittent-vent	not classified
LB06-PC05	On/Off	intermittent-vent	not classified
LB06-PC11	On/Off	intermittent-vent	not classified
LB07-PC01	On/Off	intermittent-vent	not classified
LB07-PC02	On/Off	continuous bleed *	not classified
LB07-PC03	On/Off	intermittent-vent	not classified
LB07-PC04	On/Off	intermittent-vent	not classified
OF01-PC01	On/Off	intermittent-vent *	not classified
OF02-PC01	On/Off	intermittent-vent	not classified
OF03-PC01	On/Off	intermittent-vent	not classified
OF04-PC01	On/Off	intermittent-vent	not classified
OF05-PC01	On/Off	intermittent-vent	not classified
OF06-PC01	On/Off	intermittent-vent	not classified
OF07-PC01	On/Off	continuous bleed *	not classified
OF08-PC01	On/Off	intermittent-vent	not classified
OF08-PC02	On/Off	intermittent-vent	not classified
OF09-PC01	On/Off	intermittent-vent	not classified
OF10-PC01	On/Off	continuous bleed	not classified
OF11-PC01	On/Off	intermittent-vent *	not classified
OF11-PC02	Throttle	continuous bleed *	not classified
OF11-PC03	Throttle	intermittent-vent	not classified
OF11-PC04	Throttle	intermittent-vent	not classified
RB01-PC01	On/Off	intermittent-vent	not classified
RB01-PC02	On/Off	intermittent-vent	not classified
RB01-PC03	On/Off	intermittent-vent	not classified
RB01-PC04	Throttle	continuous bleed	not classified
RB01-PC05	On/Off	intermittent-vent	not classified
RB01-PC11	On/Off	intermittent-vent	not classified
RB01-PC12	On/Off	intermittent-vent	not classified
RB01-PC13	On/Off	continuous bleed	not classified
RB01-PC14	On/Off	intermittent-vent *	not classified
RB01-PC15	On/Off	intermittent-vent	not classified
RB01-PC21	Throttle	continuous bleed	not classified

Table A-6 (continued).

Device Identifier	Field Characterization of Device service	Classification based on time series ¹	Company classification into EPA categories ²
RB01-PC23	Throttle	continuous bleed	not classified
RB01-PC24	On/Off	intermittent-vent	not classified
RB01-PC25	On/Off	intermittent-vent	not classified
RB01-PC34	On/Off	continuous bleed	not classified
RB02-PC01	On/Off	intermittent-vent	not classified
RB02-PC02	On/Off	intermittent-vent	not classified
RB02-PC03	Throttle	continuous bleed *	not classified
RB02-PC05	On/Off	intermittent-vent	not classified
RB02-PC11	On/Off	intermittent-vent	not classified
RB02-PC12	On/Off	intermittent-vent	not classified
RQ01-PC01	Throttle	intermittent-vent	not classified
RQ02-PC01	Throttle	continuous bleed	not classified
RQ02-PC02	On/Off	intermittent-vent	not classified
RQ03-PC01	Throttle	continuous bleed	not classified
RQ04-PC01	Throttle	intermittent-vent	not classified
RQ05-PC01	Throttle	intermittent-vent	not classified
RQ05-PC02	On/Off	intermittent-vent	not classified
RQ05-PC03	Throttle	continuous bleed	not classified
RQ05-PC04	On/Off	intermittent-vent	not classified
RQ06-PC01	On/Off	intermittent-vent	not classified
RQ06-PC02	On/Off	intermittent-vent	not classified
RQ06-PC03	Throttle	intermittent-vent	not classified
RQ06-PC04	Throttle	intermittent-vent	not classified
RQ07-PC01	On/Off	intermittent-vent	not classified
RQ07-PC02	On/Off	intermittent-vent	not classified
RQ07-PC03	Throttle	continuous bleed	not classified
RQ07-PC04	Throttle	continuous bleed	not classified
RQ07-PC05	On/Off	intermittent-vent	not classified
VF01-PC01	On/Off	continuous bleed	not classified
VF01-PC02	On/Off	intermittent-vent	not classified
VF01-PC03	On/Off	intermittent-vent	not classified
VF01-PC04	On/Off	intermittent-vent	not classified
VF01-PC05	On/Off	continuous bleed	not classified
VF01-PC11	On/Off	intermittent-vent	not classified
VF01-PC12	On/Off	intermittent-vent	not classified
VF01-PC13	On/Off	intermittent-vent	not classified
VF01-PC14	On/Off	intermittent-vent	not classified

Table A-6 (continued).

Device Identifier	Field Characterization of Device service	Classification based on time series ¹	Company classification into EPA categories ²
VF01-PC15	On/Off	intermittent-vent	not classified
VF01-PC21	On/Off	intermittent-vent	not classified
VF01-PC22	On/Off	continuous bleed	not classified
VF01-PC23	On/Off	intermittent-vent	not classified
VF01-PC24	On/Off	intermittent-vent	not classified
VF01-PC25	On/Off	intermittent-vent	not classified
VF01-PC31	On/Off	intermittent-vent	not classified
VF01-PC32	On/Off	intermittent-vent	not classified
VF01-PC33	On/Off	intermittent-vent	not classified
VF01-PC34	On/Off	intermittent-vent	not classified
VF01-PC35	On/Off	intermittent-vent	not classified
VF01-PC41	On/Off	intermittent-vent	not classified
VF01-PC42	On/Off	intermittent-vent	not classified
VF01-PC43	On/Off	intermittent-vent	not classified
VF01-PC44	On/Off	intermittent-vent	not classified
VF01-PC45	On/Off	continuous bleed	not classified
VF01-PC51	On/Off	continuous bleed	not classified
VF01-PC52	On/Off	intermittent-vent	not classified
VF01-PC53	On/Off	intermittent-vent *	not classified
VF01-PC54	On/Off	intermittent-vent *	not classified
VF01-PC55	On/Off	intermittent-vent	not classified
VF02-PC01	On/Off	intermittent-vent	not classified
VF02-PC02	On/Off	intermittent-vent	low bleed
VF02-PC03	On/Off	intermittent-vent	low bleed
VF02-PC04	On/Off	intermittent-vent	low bleed
VF02-PC05	On/Off	intermittent-vent	low bleed
VF02-PC11	On/Off	intermittent-vent	low bleed
VF02-PC12	On/Off	intermittent-vent	low bleed
XQ01-PC01	On/Off	intermittent-vent	intermittent
XQ01-PC02	On/Off	intermittent-vent	intermittent
XQ01-PC03	Throttle	intermittent-vent	intermittent
XQ01-PC04	On/Off	continuous bleed *	high bleed
XQ02-PC01	On/Off	intermittent-vent	intermittent
XQ02-PC02	On/Off	intermittent-vent	intermittent
XQ02-PC03	Throttle	intermittent-vent	intermittent
XQ02-PC04	On/Off	continuous bleed	high bleed
XQ03-PC01	On/Off	continuous bleed	high bleed

Table A-6 (continued).

Device Identifier	Field Characterization of Device service	Classification based on time series ¹	Company classification into EPA categories ²
XQ03-PC02	On/Off	continuous bleed	high bleed
XQ03-PC03	Throttle	intermittent-vent	intermittent
XQ03-PC04	On/Off	continuous bleed	high bleed
XQ04-PC01	On/Off	intermittent-vent	intermittent
XQ04-PC02	On/Off	continuous bleed	high bleed
XQ04-PC03	Throttle	intermittent-vent	intermittent
XQ04-PC04	On/Off	continuous bleed *	intermittent
XQ05-PC01	On/Off	intermittent-vent	intermittent
XQ05-PC02	On/Off	continuous bleed	high bleed
XQ05-PC03	Throttle	intermittent-vent	intermittent
XQ05-PC04	On/Off	intermittent-vent	intermittent
XQ06-PC01	On/Off	continuous bleed	intermittent
XQ06-PC02	On/Off	continuous bleed	intermittent
XQ06-PC03	Throttle	intermittent-vent	intermittent
XQ06-PC04	On/Off	intermittent-vent	intermittent
XQ07-PC01	On/Off	intermittent-vent	intermittent
XQ07-PC02	On/Off	intermittent-vent	intermittent
XQ07-PC03	Throttle	intermittent-vent	intermittent
XQ07-PC04	On/Off	intermittent-vent	intermittent
ZW01-PC01	On/Off	continuous bleed *	not classified
ZW01-PC02	On/Off	intermittent-vent	not classified
ZW01-PC03	On/Off	intermittent-vent	not classified
ZW01-PC04	On/Off	intermittent-vent	not classified
ZW01-PC05	On/Off	intermittent-vent	not classified
ZW01-PC11	On/Off	intermittent-vent	not classified
ZW01-PC12	On/Off	intermittent-vent	not classified
ZW01-PC13	On/Off	continuous bleed *	not classified
ZW01-PC14	On/Off	intermittent-vent	not classified
ZW01-PC15	On/Off	intermittent-vent	not classified
ZW01-PC21	On/Off	intermittent-vent	not classified
ZW01-PC22	On/Off	intermittent-vent	not classified
ZW01-PC23	On/Off	intermittent-vent	not classified
ZW01-PC24	On/Off	intermittent-vent	not classified
ZW01-PC25	On/Off	intermittent-vent	not classified
ZW01-PC31	On/Off	intermittent-vent	not classified
ZW01-PC32	On/Off	intermittent-vent	not classified
ZW01-PC33	On/Off	intermittent-vent	not classified

Table A-6 (continued).

Device Identifier	Field Characterization of Device service	Classification based on time series ¹	Company classification into EPA categories ²
ZW01-PC34	On/Off	intermittent-vent	not classified
ZW01-PC35	On/Off	intermittent-vent	not classified
ZW02-PC01	On/Off	continuous bleed	not classified
ZW02-PC02	On/Off	intermittent-vent	not classified
ZW02-PC03	On/Off	intermittent-vent	not classified
ZW02-PC04	On/Off	intermittent-vent	not classified
ZW02-PC12	On/Off	intermittent-vent	not classified
ZW03-PC01	On/Off	intermittent-vent	not classified
ZW03-PC02	On/Off	continuous bleed *	not classified
ZW03-PC03	On/Off	intermittent-vent	not classified

(1) Classification based on time series: *Intermittent-vent*: Clear actuation pattern, returning to zero between actuations. *Continuous bleed*: Non-zero, no temporal variability. A (*) Indicates that the study team made classification based on best judgment due to ambiguity from the time series.

(2) *Intermittent High bleed*, or *Low bleed*, for sites where all devices had the same classification (non-ambiguous). *Not classified* for sites where companies reported more than one device type,

Table A-7. For each device, the manufacturer and model are classified into blinded bins. The table shows measured tubing diameter and tubing length (from the controller to the valve it controls), supply gas pressure, as well as gas hydrocarbon composition for the well each device is servicing.

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
AA01-PC04	Q01	29	0.25	94	81.85	8.72	3.86	2.47
AA01-PC05	Q01	30	0.25	94	81.85	8.72	3.86	2.47
AA01-PC06	Q01	0	0.25	94	81.42	8.79	4.02	2.69
AA01-PC07	Q01	0	0.25	94	81.42	8.79	4.02	2.69
AA01-PC08	Q01	28	0.25	94	78.32	8.87	4.44	3.05
AA01-PC09	Q01	30	0.25	94	78.32	8.87	4.44	3.05
AA01-PC11	I01	34	0.25	15	81.85	8.72	3.86	2.47
AA02-PC04	H01	11	0.25	6	80.53	8.79	4.10	2.74
AA02-PC05	L01	29	0.25	32	80.53	8.79	4.10	2.74
AA02-PC06	L01	29	0.25	39	80.53	8.79	4.10	2.74
AA02-PC07	L01	29	0.25	41	80.53	8.79	4.10	2.74
AA02-PC08	I01	60	0.375	23	80.53	8.79	4.10	2.74
AA02-PC09	L01	29	0.25	19	80.53	8.79	4.10	2.74
AP01-PC01	N01	30	0.375	180	66.43	14.72	10.54	5.88
AP01-PC02	N01	30	0.375	50	66.43	14.72	10.54	5.88
AP01-PC03	N01	30	0.375	50	66.43	14.72	10.54	5.88
AP01-PC04	N01	30	0.375	60	66.43	14.72	10.54	5.88
AP01-PC05	N01	30	0.375	60	66.43	14.72	10.54	5.88
AP01-PC12	I02	11	0.375	25	66.43	14.72	10.54	5.88
AP02-PC01	I02	22	0.375	90	68.85	14.18	9.90	4.86
AP02-PC02	I02	20	0.375	90	70.63	13.28	9.05	4.85
AP02-PC03	I02	24	0.375	75	72.66	12.24	8.36	4.40

Table A-7 (continued)

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
AP02-PC04	I02	55	0.375	45	70.05	12.41	9.96	4.87
AP03-PC01	I02	30	0.375	55	64.46	15.11	11.39	6.86
AP03-PC02	I02	30	0.375	50	68.26	13.28	10.21	6.25
AP04-PC01	I02	32	0.375	25	65.35	13.15	11.62	7.04
AP04-PC02	I02	23	0.375	40	66.28	14.46	9.79	6.61
AP04-PC03	I02	32	0.375	65	62.75	13.93	12.76	7.82
AP04-PC04	I02	30	0.375	55	59.63	16.87	14.22	7.25
AP05-PC01	I02	34	0.375	25	70.86	14.03	7.87	5.14
AP05-PC02	I02	32	0.375	70	70.86	14.03	7.87	5.14
CW01-PC01	F01	20	0.375	45	81.26	7.76	3.90	2.87
CW01-PC02	B01	20	0.375	130	81.26	7.76	3.90	2.87
CW01-PC03	Q02	22	0.375	30	81.26	7.76	3.90	2.87
CW01-PC04	Q02	18	0.375	30	81.26	7.76	3.90	2.87
CW01-PC05	I05	20	0.375	28	81.26	7.76	3.90	2.87
CW01-PC11	F01	25	0.375	45	81.58	8.25	3.75	2.71
CW01-PC12	B01	25	0.375	130	81.58	8.25	3.75	2.71
CW01-PC13	Q02	25	0.375	30	81.58	8.25	3.75	2.71
CW01-PC14	Q02	30	0.375	30	81.58	8.25	3.75	2.71
CW01-PC15	I05	25	0.375	28	81.58	8.25	3.75	2.71
CW01-PC21	F01	18	0.375	45	84.18	7.69	2.84	1.85
CW01-PC22	B01	18	0.375	130	84.18	7.69	2.84	1.85
CW01-PC23	Q02	18	0.375	30	84.18	7.69	2.84	1.85
CW01-PC24	Q02	18	0.375	30	84.18	7.69	2.84	1.85
CW01-PC25	I05	18	0.375	28	84.18	7.69	2.84	1.85

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
CW01-PC31	F01	20	0.375	45	81.69	7.79	3.77	2.60
CW01-PC32	B01	20	0.375	130	81.69	7.79	3.77	2.60
CW01-PC33	Q02	22	0.375	30	81.69	7.79	3.77	2.60
CW01-PC34	Q02	22	0.375	30	81.69	7.79	3.77	2.60
CW01-PC35	I05	22	0.375	28	81.69	7.79	3.77	2.60
CW02-PC01	F01	20	0.375	45	81.80	8.12	4.39	2.63
CW02-PC02	B01	20	0.375	130	81.80	8.12	4.39	2.63
CW02-PC03	Q02	20	0.375	30	81.80	8.12	4.39	2.63
CW02-PC04	Q02	19	0.375	30	81.80	8.12	4.39	2.63
CW02-PC05	I05	20	0.375	28	81.80	8.12	4.39	2.63
CW02-PC11	F01	15	0.375	45	80.64	7.81	4.41	3.51
CW02-PC12	B01	15	0.375	130	80.64	7.81	4.41	3.51
CW02-PC13	Q02	16	0.375	30	80.64	7.81	4.41	3.51
CW02-PC14	Q02	16	0.375	30	80.64	7.81	4.41	3.51
CW02-PC15	I05	16	0.375	28	80.64	7.81	4.41	3.51
CW02-PC21	F01	20	0.375	45	82.36	7.66	3.78	2.60
CW02-PC22	B01	20	0.375	130	82.36	7.66	3.78	2.60
CW02-PC23	Q02	22	0.375	30	82.36	7.66	3.78	2.60
CW02-PC24	Q02	22	0.375	30	82.36	7.66	3.78	2.60
CW02-PC25	I05	22	0.375	28	82.36	7.66	3.78	2.60
CW02-PC31	F01	19	0.375	30	82.02	7.89	4.01	2.48
CW02-PC32	B01	19	0.375	130	82.02	7.89	4.01	2.48
CW02-PC33	Q02	19	0.375	30	82.02	7.89	4.01	2.48
CW02-PC34	Q02	20	0.375	30	82.02	7.89	4.01	2.48

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
CW02-PC35	I05	19	0.375	28	82.02	7.89	4.01	2.48
CZ01-PC01	M01	32.5	0.375	40	90.81	3.25	0.99	2.26
CZ01-PC02	E01	20	0.375	11	90.81	3.25	0.99	2.26
CZ02-PC01	J01	17	0.25	54	91.13	2.89	0.91	2.26
CZ02-PC02	E01	20	0.25	32	91.13	2.89	0.91	2.26
CZ03-PC01	A01	27	0.5	37	93.33	2.04	0.51	1.26
CZ03-PC02	E01	20	0.375	13	93.33	2.04	0.51	1.26
CZ04-PC01	M01	11	0.375	43	91.72	2.82	0.79	1.89
CZ04-PC02	E01	20	0.375	10	91.72	2.82	0.79	1.89
CZ05-PC01	A01	29	0.375	45	93.76	1.87	0.46	1.14
CZ05-PC02	E01	20	0.375	19	93.76	1.87	0.46	1.14
CZ05-PC03	A01	29	0.375	18	93.76	1.87	0.46	1.14
CZ06-PC01	G01	6	0.25	73	94.07	1.84	0.42	0.92
CZ06-PC02	E01	20	0.375	17	94.07	1.84	0.42	0.92
CZ06-PC03	G01	6	0.25	39	94.07	1.84	0.42	0.92
CZ07-PC01	A01	20	0.375	38	93.29	2.10	0.54	1.20
CZ07-PC02	E01	27.5	0.375	27	93.29	2.10	0.54	1.20
CZ08-PC01	M01	11	0.375	77	96.14	0.76	0.11	0.20
CZ08-PC02	M01	19	0.375	77	97.58	0.78	0.08	0.15
CZ09-PC01	M01	19	0.375	52	96.07	0.70	0.06	0.10
CZ09-PC02	M01	9	0.375	40	96.07	0.70	0.06	0.10
CZ10-PC01	M01	13	0.375	40	96.10	0.69	0.09	0.19
CZ11-PC01	M01	23	0.375	43	95.81	0.86	0.11	0.20
DL01-PC01	Q02	15	0.375	40	90.57	4.93	1.37	1.14

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
DL01-PC02	Q02	12	0.375	40	90.57	4.93	1.37	1.14
DL01-PC03	I05	30	0.375	90	90.57	4.93	1.37	1.14
DL01-PC04	F01	27	0.375	90	90.57	4.93	1.37	1.14
DL01-PC05	I05	25	0.375	90	90.84	4.92	1.43	1.16
DL01-PC11	M01	30	0.375	25	90.84	4.92	1.43	1.16
DL01-PC12	F01	27	0.375	160	90.84	4.92	1.43	1.16
DL01-PC13	M01	35	0.375	25	90.10	5.08	1.39	1.08
DL01-PC14	F01	30	0.375	160	90.10	5.08	1.39	1.08
DL01-PC15	M01	35	0.375	25	91.00	4.16	1.04	0.88
DL01-PC21	F01	35	0.375	160	91.00	4.16	1.04	0.88
DL01-PC22	M01	30	0.375	25	90.65	4.82	1.32	1.12
DL01-PC23	F01	30	0.375	160	90.65	4.82	1.32	1.12
DL01-PC24	M01	38	0.375	25	89.35	5.19	1.39	1.18
DL01-PC25	F01	28	0.375	160	89.35	5.19	1.39	1.18
DL01-PC31	I05	25	0.375	90	90.84	4.92	1.43	1.16
DL01-PC32	M01	28	0.375	25	90.84	4.92	1.43	1.16
DL01-PC33	F01	28	0.375	160	90.84	4.92	1.43	1.16
DL01-PC34	M01	26	0.375	25	90.88	4.82	1.35	1.16
DL01-PC35	F01	25	0.375	160	90.88	4.82	1.35	1.16
DL01-PC41	M01	28	0.375	25	89.68	5.11	1.38	1.11
DL01-PC42	F01	28	0.375	160	89.68	5.11	1.38	1.11
DL01-PC43	M01	29	0.375	25	89.35	5.30	1.44	1.03
DL01-PC44	F01	28	0.375	160	89.35	5.30	1.44	1.03
DL01-PC45	I05	25	0.375	45	89.35	5.30	1.44	1.03

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
DL02-PC01	M03	28	0.375	25	90.36	4.93	1.30	1.03
DL02-PC02	F01	28	0.375	160	90.36	4.93	1.30	1.03
DL02-PC03	M03	22	0.375	25	90.51	5.03	1.37	1.09
DL02-PC04	F01	25	0.375	160	90.51	5.03	1.37	1.09
DL02-PC05	M03	26	0.375	25	90.51	5.03	1.37	1.09
DL02-PC11	F01	25	0.375	160	90.51	5.03	1.37	1.09
DL02-PC12	M03	24	0.375	25	90.39	4.99	1.34	1.03
DL02-PC13	F01	25	0.375	160	90.39	4.99	1.34	1.03
DL02-PC14	M03	26	0.375	25	90.47	5.19	1.47	1.11
DL02-PC15	F01	25	0.375	160	90.47	5.19	1.47	1.11
DL02-PC21	M03	26	0.375	25	90.20	4.98	1.30	1.04
DL02-PC22	F01	27	0.375	160	90.20	4.98	1.30	1.04
DL02-PC23	M03	26	0.375	25	90.00	5.03	1.35	0.88
DL02-PC24	F01	26	0.375	160	90.00	5.03	1.35	0.88
DL02-PC25	M03	26	0.375	25	90.21	5.10	1.40	1.04
DL02-PC31	F01	27	0.375	160	90.21	5.10	1.40	1.04
DL02-PC32	M03	26	0.375	25	89.80	5.15	1.31	0.97
DL02-PC33	F01	26	0.375	160	89.80	5.15	1.31	0.97
DL02-PC34	M03	28	0.375	25	90.38	5.08	1.38	0.84
DL02-PC35	F01	28	0.375	160	90.38	5.08	1.38	0.84
DL02-PC41	M03	30	0.375	25	90.06	4.97	1.28	0.82
DL02-PC42	F01	28	0.375	160	90.06	4.97	1.28	0.82
DL02-PC43	M03	28	0.375	25	90.27	4.78	1.20	1.50
DL02-PC44	F01	28	0.375	160	90.27	4.78	1.20	1.50

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
DL02-PC45	M03	28	0.375	25	90.38	4.96	1.32	0.97
DL02-PC51	F01	26	0.375	160	90.38	4.96	1.32	0.97
DL02-PC52	M03	28	0.375	25	89.67	5.07	1.17	0.81
DL02-PC53	F01	28	0.375	160	89.67	5.07	1.17	0.81
DL02-PC54	I05	25	0.375	90	90.20	4.98	1.30	1.04
DL02-PC55	I05	28	0.375	90	90.20	4.98	1.30	1.04
DL02-PC61	I05	25	0.375	90	90.20	4.98	1.30	1.04
DL02-PC62	I05	25	0.375	45	90.20	4.98	1.30	1.04
GZ01-PC01	J01	23	0.375	47	97.40	0.52	0.11	0.21
GZ01-PC02	J01	22	0.375	47	97.40	0.52	0.11	0.21
GZ01-PC03	J01	26	0.375	184	97.42	0.52	0.11	0.15
GZ01-PC04	J01	26	0.375	47	97.42	0.52	0.11	0.15
GZ01-PC05	J01	20	0.375	184	97.32	0.52	0.11	0.25
GZ01-PC11	J01	23	0.375	47	97.32	0.52	0.11	0.25
GZ01-PC12	J01	25	0.375	184	97.29	0.55	0.12	0.28
GZ01-PC13	J01	23	0.375	47	97.29	0.55	0.12	0.28
GZ01-PC14	K01	30	0.375	NA	97.40	0.52	0.11	0.21
GZ01-PC15	K01	30	0.375	NA	97.42	0.52	0.11	0.15
GZ01-PC21	K01	30	0.375	NA	97.32	0.52	0.11	0.25
GZ01-PC22	K01	30	0.375	NA	97.29	0.55	0.12	0.28
GZ02-PC01	M01	29	0.375	127	97.35	0.51	0.10	0.21
GZ02-PC02	M01	28	0.375	35	97.35	0.51	0.10	0.21
GZ02-PC03	M01	30	0.375	127	97.42	0.51	0.09	0.12
GZ02-PC04	M01	31	0.375	35	97.42	0.51	0.09	0.12

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
GZ02-PC05	H01	NA	0.375	NA	97.42	0.51	0.09	0.12
GZ02-PC11	J01	30	0.375	110	97.42	0.51	0.09	0.12
GZ02-PC12	J01	28	0.375	120	97.42	0.51	0.09	0.12
GZ02-PC13	J01	30	0.375	135	97.42	0.51	0.09	0.12
GZ02-PC14	I01	42	0.375	36	97.42	0.51	0.09	0.12
GZ02-PC15	M01	NA	0.375	20	97.42	0.51	0.09	0.12
GZ02-PC21	M01	20	0.25	120	97.42	0.51	0.09	0.12
GZ02-PC22	K01	3	0.375	3600	97.42	0.51	0.09	0.12
GZ02-PC23	K01	3	0.375	3600	97.35	0.51	0.10	0.21
GZ02-PC24	I01	300	0.375	25	97.42	0.51	0.09	0.12
GZ03-PC01	L01	30	0.25	26	97.16	0.33	0.03	0.08
GZ03-PC02	L01	30	0.25	26	97.16	0.33	0.03	0.08
GZ03-PC03	L01	30	0.25	26	97.16	0.33	0.03	0.08
GZ03-PC04	H01	NA	0.25	8	97.16	0.33	0.03	0.08
GZ03-PC05	I01	32	0.375	36	97.16	0.33	0.03	0.08
GZ03-PC11	K01	37	0.375	5400	97.16	0.33	0.03	0.08
GZ03-PC12	K01	37	0.375	5400	97.16	0.33	0.03	0.08
GZ03-PC13	M01	26	0.375	93	97.16	0.33	0.03	0.08
GZ03-PC14	M01	27	0.375	70	97.16	0.33	0.03	0.08
GZ03-PC15	M01	25	0.375	55	97.16	0.33	0.03	0.08
GZ03-PC21	M01	27	0.375	84	97.16	0.33	0.03	0.08
GZ03-PC22	M01	7	0.375	47	97.16	0.33	0.03	0.08
GZ03-PC23	M01	32	0.375	29	97.36	0.30	0.03	0.04
GZ03-PC24	M01	1	0.375	47	97.36	0.30	0.03	0.04

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
GZ03-PC25	K01	31	0.375	66	97.16	0.33	0.03	0.08
GZ03-PC31	M04	25	0.375	95	97.16	0.33	0.03	0.08
GZ03-PC32	M03	26	0.25	28	97.16	0.33	0.03	0.08
GZ03-PC33	M03	29	0.25	86	97.16	0.33	0.03	0.08
GZ03-PC34	I05	NA	0.375	40	97.16	0.33	0.03	0.08
GZ03-PC35	I05	29	0.375	31	97.16	0.33	0.03	0.08
GZ03-PC41	H01	32	0.25	8	97.16	0.33	0.03	0.08
GZ03-PC42	I01	36	0.375	20	97.16	0.33	0.03	0.08
GZ04-PC01	M01	27	0.375	94	90.83	0.22	0.01	0.07
GZ04-PC02	M01	25	0.6	32	90.83	0.22	0.01	0.07
GZ04-PC03	M01	30	0.375	25	97.44	0.23	0.02	0.02
GZ04-PC04	M01	30	0.375	94	97.44	0.23	0.02	0.02
GZ04-PC05	K01	30	0.375	5400	90.83	0.22	0.01	0.07
GZ04-PC11	K01	32	0.375	3600	97.44	0.23	0.02	0.02
LB01-PC01	M03	30	0.375	400	80.52	10.62	3.62	3.61
LB01-PC02	M03	30	0.375	292	80.52	10.62	3.62	3.61
LB02-PC01	M01	30	0.375	85	80.41	10.77	3.56	3.48
LB02-PC02	M01	30	0.375	80	80.41	10.77	3.56	3.48
LB02-PC03	M01	30	0.375	80	80.41	10.77	3.56	3.48
LB02-PC04	M01	30	0.375	80	80.41	10.77	3.56	3.48
LB03-PC01	M01	30	0.375	73	79.46	10.95	4.15	4.05
LB03-PC02	M01	30	0.375	48	79.46	10.95	4.15	4.05
LB04-PC01	M03	30	0.375	90	79.15	11.16	4.26	3.94
LB04-PC02	M03	30	0.375	85	79.15	11.16	4.26	3.94

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
LB04-PC03	M03	30	0.375	93	78.53	11.43	4.43	4.45
LB04-PC04	M03	30	0.375	83	78.53	11.43	4.43	4.45
LB05-PC01	M03	30	0.375	390	87.73	3.07	0.81	0.99
LB05-PC02	M03	30	0.375	328	87.73	3.07	0.81	0.99
LB05-PC03	M03	25	0.375	387	87.73	3.07	0.81	0.99
LB05-PC04	M03	25	0.375	352	87.73	3.07	0.81	0.99
LB06-PC01	M03	32	0.375	408	81.15	11.16	3.75	2.65
LB06-PC02	M03	32	0.375	301	81.15	11.16	3.75	2.65
LB06-PC03	M03	32	0.375	427	80.67	11.27	3.88	2.92
LB06-PC04	M03	28	0.375	300	80.67	11.27	3.88	2.92
LB06-PC05	M03	30	0.375	427	81.44	11.10	3.66	2.50
LB06-PC11	M03	30	0.375	300	81.44	11.10	3.66	2.50
LB07-PC01	M03	24	0.375	85	79.04	11.45	4.24	3.97
LB07-PC02	M03	24	0.375	85	79.04	11.45	4.24	3.97
LB07-PC03	J01	28	0.375	85	79.31	11.34	4.14	3.99
LB07-PC04	M03	28	0.375	85	79.31	11.34	4.14	3.99
OF01-PC01	I03	14	0.25	34	96.34	2.96	0.43	0.22
OF02-PC01	I03	20	0.25	35	96.34	2.96	0.43	0.22
OF03-PC01	I03	30	0.25	38	96.34	2.96	0.43	0.22
OF04-PC01	I03	18	0.25	50	96.34	2.96	0.43	0.22
OF05-PC01	I03	30	0.25	39	96.34	2.96	0.43	0.22
OF06-PC01	I03	16	0.25	39	96.34	2.96	0.43	0.22
OF07-PC01	I03	22	0.375	88	96.34	2.96	0.43	0.22
OF08-PC01	I03	27	0.25	110	96.34	2.96	0.43	0.22

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
OF08-PC02	E01	NA	0.25	20	96.34	2.96	0.43	0.22
OF09-PC01	I03	14	0.25	25	96.34	2.96	0.43	0.22
OF10-PC01	I03	23	0.25	20	96.34	2.96	0.43	0.22
OF11-PC01	I03	29	0.25	23	96.34	2.96	0.43	0.22
OF11-PC02	M01	40	0.375	120	96.34	2.96	0.43	0.22
OF11-PC03	M01	40	0.375	92	96.34	2.96	0.43	0.22
OF11-PC04	I01	40	0.375	18	96.34	2.96	0.43	0.22
RB01-PC01	M03	10	0.25	1	82.19	8.47	3.85	3.33
RB01-PC02	M03	32	0.375	140	82.19	8.47	3.85	3.33
RB01-PC03	M03	32	0.375	140	82.19	8.47	3.85	3.33
RB01-PC04	M05	30	0.375	15	82.19	8.47	3.85	3.33
RB01-PC05	M01	35	0.375	45	82.19	8.47	3.85	3.33
RB01-PC11	M01	32	0.375	70	82.19	8.47	3.85	3.33
RB01-PC12	M01	32	0.375	180	82.19	8.47	3.85	3.33
RB01-PC13	M01	25	0.375	100	82.19	8.47	3.85	3.33
RB01-PC14	M01	19	0.375	60	82.19	8.47	3.85	3.33
RB01-PC15	M01	32	0.375	NA	82.19	8.47	3.85	3.33
RB01-PC21	I05	10	0.25	100	82.19	8.47	3.85	3.33
RB01-PC23	M05	30	0.375	30	82.19	8.47	3.85	3.33
RB01-PC24	M01	32	0.375	30	82.19	8.47	3.85	3.33
RB01-PC25	H01	52	0.375	1	82.19	8.47	3.85	3.33
RB01-PC34	M01	32	0.375	140	82.19	8.47	3.85	3.33
RB02-PC01	M01	32	0.375	75	72.35	14.60	7.40	3.44
RB02-PC02	M01	29	0.25	25	72.35	14.60	7.40	3.44

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
RB02-PC03	I05	14	0.25	35	72.35	14.60	7.40	3.44
RB02-PC05	M01	30	0.25	100	72.35	14.60	7.40	3.44
RB02-PC11	M01	30	0.25	100	72.35	14.60	7.40	3.44
RB02-PC12	M01	32	0.25	NA	72.35	14.60	7.40	3.44
RQ01-PC01	Q02	24	0.25	70	86.88	6.48	2.83	2.48
RQ02-PC01	Q02	23	0.25	32	86.22	6.43	2.86	3.21
RQ02-PC02	O01	30	0.375	102	86.22	6.43	2.86	3.21
RQ03-PC01	Q02	22	0.375	40	85.61	6.43	2.84	3.80
RQ04-PC01	Q02	20	0.25	28	86.24	6.45	2.84	3.16
RQ05-PC01	Q02	21	0.375	34	88.27	6.12	2.86	1.52
RQ05-PC02	O01	30	0.375	34	88.27	6.12	2.86	1.52
RQ05-PC03	G01	22	0.375	38	87.71	6.11	3.20	1.76
RQ05-PC04	O01	30	0.375	0	87.71	6.11	3.20	1.76
RQ06-PC01	I05	8	0.25	32	84.16	8.24	3.97	1.61
RQ06-PC02	I05	8	0.25	47	84.16	8.24	3.97	1.61
RQ06-PC03	M03	27	0.25	38	84.16	8.24	3.97	1.61
RQ06-PC04	M03	29	0.25	27	84.16	8.24	3.97	1.61
RQ07-PC01	I05	18	0.25	65	77.73	9.13	6.59	4.32
RQ07-PC02	I05	18	0.25	55	77.73	9.13	6.59	4.32
RQ07-PC03	G01	22	0.25	36	77.73	9.13	6.59	4.32
RQ07-PC04	G01	21	0.25	25	77.73	9.13	6.59	4.32
RQ07-PC05	P01	25	0.375	0	77.73	9.13	6.59	4.32
VF01-PC01	F01	95	0.375	140	97.49	2.02	0.06	0.00
VF01-PC02	M02	40	0.375	90	97.49	2.02	0.06	0.00

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
VF01-PC03	I04	21	0.25	27	97.49	2.02	0.06	0.00
VF01-PC04	I04	21	0.25	40	97.49	2.02	0.06	0.00
VF01-PC05	F01	61	0.25	134	97.63	1.95	0.05	0.00
VF01-PC11	M02	38	0.375	90	97.63	1.95	0.05	0.00
VF01-PC12	I04	23	0.25	15	97.63	1.95	0.05	0.00
VF01-PC13	I04	22	0.25	41	97.63	1.95	0.05	0.00
VF01-PC14	F01	50	0.375	95	97.61	2.01	0.06	0.00
VF01-PC15	M02	35	0.375	90	97.61	2.01	0.06	0.00
VF01-PC21	I04	7	0.25	26	97.61	2.01	0.06	0.00
VF01-PC22	I04	7	0.25	45	97.61	2.01	0.06	0.00
VF01-PC23	F01	86	0.375	110	97.74	1.82	0.04	0.00
VF01-PC24	M02	34	0.375	100	97.74	1.82	0.04	0.00
VF01-PC25	I04	15	0.375	32	97.74	1.82	0.04	0.00
VF01-PC31	I04	15	0.375	45	97.74	1.82	0.04	0.00
VF01-PC32	F01	25	0.375	98	97.63	2.01	0.05	0.00
VF01-PC33	M02	40	0.375	90	97.63	2.01	0.05	0.00
VF01-PC34	I04	14	0.25	25	97.63	2.01	0.05	0.00
VF01-PC35	I04	14	0.25	40	97.63	2.01	0.05	0.00
VF01-PC41	F01	89	0.375	84	97.68	1.89	0.05	0.00
VF01-PC42	M02	36	0.375	85	97.68	1.89	0.05	0.00
VF01-PC43	I04	17	0.25	26	97.68	1.89	0.05	0.00
VF01-PC44	I04	17	0.25	40	97.68	1.89	0.05	0.00
VF01-PC45	C01	27	0.375	28	97.49	2.02	0.06	0.00
VF01-PC51	C01	18	0.375	28	97.63	1.95	0.05	0.00

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
VF01-PC52	C01	30	0.375	28	97.61	2.01	0.06	0.00
VF01-PC53	C01	25	0.375	28	97.74	1.82	0.04	0.00
VF01-PC54	C01	25	0.375	28	97.63	2.01	0.05	0.00
VF01-PC55	C01	30	0.375	28	97.68	1.89	0.05	0.00
VF02-PC01	F03	25	0.375	110	96.90	2.66	0.10	0.00
VF02-PC02	F03	30	0.375	110	96.94	2.59	0.09	0.00
VF02-PC03	F03	30	0.375	110	96.81	2.68	0.09	0.00
VF02-PC04	F03	30	0.375	110	96.98	2.56	0.08	0.00
VF02-PC05	F03	28	0.375	110	97.02	2.66	0.09	0.00
VF02-PC11	F03	31	0.375	110	96.93	2.61	0.09	0.00
VF02-PC12	I05	31	0.375	331	96.98	2.56	0.08	0.00
XQ01-PC01	Q02	27	0.25	30	88.21	6.64	1.98	1.46
XQ01-PC02	Q02	24	0.25	30	88.21	6.64	1.98	1.46
XQ01-PC03	I05	15	0.25	40	88.21	6.64	1.98	1.46
XQ01-PC04	F02	15	0.375	25	88.21	6.64	1.98	1.46
XQ02-PC01	G01	28	0.25	35	86.45	7.49	2.61	1.90
XQ02-PC02	G01	28	0.25	35	86.45	7.49	2.61	1.90
XQ02-PC03	I05	18	0.25	45	86.45	7.49	2.61	1.90
XQ02-PC04	F02	18	0.375	15	86.45	7.49	2.61	1.90
XQ03-PC01	Q02	23	0.25	30	90.85	5.17	1.28	1.04
XQ03-PC02	Q02	23	0.25	30	90.85	5.17	1.28	1.04
XQ03-PC03	I05	22	0.25	20	90.85	5.17	1.28	1.04
XQ03-PC04	F02	22	0.375	35	90.85	5.17	1.28	1.04
XQ04-PC01	Q02	25	0.25	38	89.49	5.33	2.32	2.27

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
XQ04-PC02	Q02	28	0.25	38	89.49	5.33	2.32	2.27
XQ04-PC03	I05	15	0.25	15	89.49	5.33	2.32	2.27
XQ04-PC04	D01	15	0.375	10	89.49	5.33	2.32	2.27
XQ05-PC01	Q02	25	0.25	35	90.16	5.38	2.11	1.75
XQ05-PC02	Q02	27	0.25	35	90.16	5.38	2.11	1.75
XQ05-PC03	I05	17	0.25	12	90.16	5.38	2.11	1.75
XQ05-PC04	D01	17	0.375	10	90.16	5.38	2.11	1.75
XQ06-PC01	G01	28	0.25	38	89.14	5.48	2.41	2.45
XQ06-PC02	G01	28	0.25	38	89.14	5.48	2.41	2.45
XQ06-PC03	I05	28	0.25	30	89.14	5.48	2.41	2.45
XQ06-PC04	D01	28	0.375	10	89.14	5.48	2.41	2.45
XQ07-PC01	G01	60	0.25	38	91.62	4.41	1.72	1.59
XQ07-PC02	G01	60	0.25	38	91.62	4.41	1.72	1.59
XQ07-PC03	I05	60	0.25	30	91.62	4.41	1.72	1.59
XQ07-PC04	D01	60	0.375	10	91.62	4.41	1.72	1.59
ZW01-PC01	Q02	28	0.375	42	90.61	5.09	2.00	1.62
ZW01-PC02	Q02	27	0.375	52	90.61	5.09	2.00	1.62
ZW01-PC03	I05	14	0.375	NA	90.61	5.09	2.00	1.62
ZW01-PC04	I05	14	0.375	NA	90.61	5.09	2.00	1.62
ZW01-PC05	I05	12	0.375	NA	90.61	5.09	2.00	1.62
ZW01-PC11	I05	12	0.375	NA	90.61	5.09	2.00	1.62
ZW01-PC12	Q02	25	0.375	42	90.87	5.24	1.80	1.25
ZW01-PC13	Q02	25	0.375	52	90.87	5.24	1.80	1.25
ZW01-PC14	Q02	25	0.375	42	90.87	5.24	1.80	1.25

Table A-7 (continued).

Device Identifier	Man., Model	Supply gas pressure (psig)	Tubing Diameter (in)	Tubing Length (in)	% of C1 in gas (%mol) ¹	% of C2 in gas (%mol) ¹	% of C3 in gas (%mol) ¹	% of C4+ in gas (%mol) ¹
ZW01-PC15	Q02	25	0.375	52	90.87	5.24	1.80	1.25
ZW01-PC21	Q02	40	0.375	22	90.87	5.24	1.80	1.25
ZW01-PC22	I05	15	0.375	NA	90.87	5.24	1.80	1.25
ZW01-PC23	I05	15	0.375	NA	90.03	5.89	2.03	1.25
ZW01-PC24	I05	15	0.375	NA	90.03	5.89	2.03	1.25
ZW01-PC25	I05	15	0.375	NA	90.03	5.89	2.03	1.25
ZW01-PC31	B01	22	0.375	60	90.03	5.89	2.03	1.25
ZW01-PC32	I05	15	0.375	NA	90.03	5.89	2.03	1.25
ZW01-PC33	I05	15	0.375	NA	90.03	5.89	2.03	1.25
ZW01-PC34	I05	15	0.375	NA	90.03	5.89	2.03	1.25
ZW01-PC35	I05	15	0.375	NA	90.03	5.89	2.03	1.25
ZW02-PC01	Q02	25	0.375	30	89.65	5.68	2.22	1.59
ZW02-PC02	Q02	25	0.375	35	89.65	5.68	2.22	1.59
ZW02-PC03	I05	60	0.375	NA	89.65	5.68	2.22	1.59
ZW02-PC04	I05	60	0.375	NA	89.65	5.68	2.22	1.59
ZW02-PC12	B01	25	0.375	52	89.65	5.68	2.22	1.59
ZW03-PC01	Q02	25	0.375	26	91.22	4.94	1.81	1.25
ZW03-PC02	Q02	25	0.375	40	91.22	4.94	1.81	1.25
ZW03-PC03	F01	25	0.375	50	91.22	4.94	1.81	1.20

(1) Total may not sum to 100% due to nitrogen, CO₂, and other species in the gas.

Table A-8. Average measured emission rates for each device (whole gas and methane).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
AA01-PC04	7.595	not measured	7.595	6.216	c
AA01-PC05	0	not measured	0.000	0.000	
AA01-PC06	1.602	not measured	1.602	1.304	c
AA01-PC07	0.013	not measured	0.013	0.011	
AA01-PC08	1.741	not measured	1.741	1.363	c
AA01-PC09	0.035	not measured	0.035	0.027	c
AA01-PC11	0.279	not measured	0.279	0.228	c
AA02-PC04	13.597	not measured	13.597	10.950	
AA02-PC05	2.514	not measured	2.514	2.025	c
AA02-PC06	4.926	not measured	4.926	3.967	
AA02-PC07	1.186	not measured	1.186	0.955	
AA02-PC08	111.413	not measured	111.413	89.718	
AA02-PC09	7.766	not measured	7.766	6.254	
AP01-PC01	1.027	not measured	1.027	0.682	c
AP01-PC02	0.001	not measured	0.001	0.000	c
AP01-PC03	0.036	not measured	0.036	0.024	
AP01-PC04	0.003	not measured	0.003	0.002	
AP01-PC05	0.001	not measured	0.001	0.001	
AP01-PC12	25.559	not measured	25.559	16.980	
AP02-PC01	0.028	not measured	0.028	0.020	
AP02-PC02	0.112	not measured	0.112	0.079	c
AP02-PC03	0.002	0.000	0.002	0.002	

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
AP02-PC04	0.032	not measured	0.032	0.022	c
AP03-PC01	0	not measured	0.000	0.000	
AP03-PC02	0	not measured	0.000	0.000	
AP04-PC01	0.004	not measured	0.004	0.002	
AP04-PC02	0.111	0.000	0.111	0.074	c
AP04-PC03	0.036	0.007	0.036	0.023	c
AP04-PC04	0.001	not measured	0.001	0.001	
AP05-PC01	0.169	0.197	0.169	0.120	c
AP05-PC02	0	not measured	0.000	0.000	
CW01-PC01	not measured	0.000	0.000	0.000	
CW01-PC02	0	not measured	0.000	0.000	
CW01-PC03	0	not measured	0.000	0.000	
CW01-PC04	0	not measured	0.000	0.000	a
CW01-PC05	0.008	not measured	0.008	0.007	a
CW01-PC11	0.022	not measured	0.022	0.018	c
CW01-PC12	0.034	not measured	0.034	0.028	a
CW01-PC13	0.235	not measured	0.235	0.192	
CW01-PC14	0.031	not measured	0.031	0.025	a
CW01-PC15	0	not measured	0.000	0.000	
CW01-PC21	0	0.002	0.000	0.000	
CW01-PC22	0	not measured	0.000	0.000	
CW01-PC23	1.377	not measured	1.377	1.159	c

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
CW01-PC24	0	not measured	0.000	0.000	a
CW01-PC25	0.003	not measured	0.003	0.003	a
CW01-PC31	0	not measured	0.000	0.000	a
CW01-PC32	0.102	not measured	0.102	0.083	b, c
CW01-PC33	0	not measured	0.000	0.000	
CW01-PC34	0	not measured	0.000	0.000	a
CW01-PC35	0.622	not measured	0.622	0.508	a, b
CW02-PC01	not measured	0.082	0.082	0.067	
CW02-PC02	0	not measured	0.000	0.000	
CW02-PC03	0	not measured	0.000	0.000	
CW02-PC04	0	not measured	0.000	0.000	a
CW02-PC05	0.004	not measured	0.004	0.003	a
CW02-PC11	not measured	0.000	0.000	0.000	
CW02-PC12	0	not measured	0.000	0.000	
CW02-PC13	0	not measured	0.000	0.000	
CW02-PC14	0	not measured	0.000	0.000	a
CW02-PC15	0.002	not measured	0.002	0.002	a
CW02-PC21	not measured	0.000	0.000	0.000	
CW02-PC22	0.003	not measured	0.003	0.003	a
CW02-PC23	0	not measured	0.000	0.000	a
CW02-PC24	0	0.142	0.000	0.000	
CW02-PC25	0	not measured	0.000	0.000	

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
CW02-PC31	not measured	0.000	0.000	0.000	
CW02-PC32	0.529	not measured	0.529	0.434	
CW02-PC33	7.821	not measured	7.821	6.415	
CW02-PC34	0	not measured	0.000	0.000	a
CW02-PC35	0.010	not measured	0.010	0.008	a
CZ01-PC01	0.243	not measured	0.243	0.221	a
CZ01-PC02	not measured	0.000	0.000	0.000	
CZ02-PC01	118.497	not measured	118.497	107.983	a
CZ02-PC02	not measured	0.580	0.580	0.528	
CZ03-PC01	0.006	not measured	0.006	0.005	a
CZ03-PC02	not measured	0.000	0.000	0.000	
CZ04-PC01	0	not measured	0.000	0.000	a
CZ04-PC02	not measured	0.000	0.000	0.000	
CZ05-PC01	40.400	not measured	40.400	37.880	a
CZ05-PC02	not measured	0.000	0.000	0.000	
CZ05-PC03	0	not measured	0.000	0.000	
CZ06-PC01	not measured	51.231	51.231	48.192	
CZ06-PC02	not measured	50.714	50.714	47.705	
CZ06-PC03	not measured	0.000	0.000	0.000	
CZ07-PC01	0.007	not measured	0.007	0.006	a
CZ07-PC02	not measured	0.000	0.000	0.000	
CZ08-PC01	60.099	not measured	60.099	57.783	a

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
CZ08-PC02	21.578	not measured	21.578	21.056	
CZ09-PC01	149.678	not measured	149.678	143.801	a
CZ09-PC02	44.871	not measured	44.871	43.109	
CZ10-PC01	43.173	not measured	43.173	41.489	a
CZ11-PC01	22.183	not measured	22.183	21.253	a
DL01-PC01	0.065	not measured	0.065	0.059	
DL01-PC02	0	not measured	0.000	0.000	a
DL01-PC03	0	not measured	0.000	0.000	a
DL01-PC04	not measured	1.461	1.461	1.323	
DL01-PC05	0	not measured	0.000	0.000	a
DL01-PC11	0	not measured	0.000	0.000	a
DL01-PC12	0.502	not measured	0.502	0.456	
DL01-PC13	0	not measured	0.000	0.000	a
DL01-PC14	0.595	not measured	0.595	0.536	
DL01-PC15	5.386	not measured	5.386	4.902	a
DL01-PC21	0.980	not measured	0.980	0.892	c
DL01-PC22	0	not measured	0.000	0.000	a
DL01-PC23	1.574	not measured	1.574	1.427	c
DL01-PC24	19.759	not measured	19.759	17.656	a
DL01-PC25	0	not measured	0.000	0.000	
DL01-PC31	0	not measured	0.000	0.000	
DL01-PC32	0	not measured	0.000	0.000	a

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
DL01-PC33	0.114	not measured	0.114	0.104	
DL01-PC34	0	not measured	0.000	0.000	a
DL01-PC35	0	not measured	0.000	0.000	
DL01-PC41	0	not measured	0.000	0.000	a
DL01-PC42	0.489	not measured	0.489	0.438	
DL01-PC43	0	not measured	0.000	0.000	a
DL01-PC44	0.502	not measured	0.502	0.449	
DL01-PC45	not measured	0.000	0.000	0.000	
DL02-PC01	0.005	not measured	0.005	0.004	a
DL02-PC02	0.465	not measured	0.465	0.420	
DL02-PC03	0.008	59.833	0.008	0.007	a
DL02-PC04	0.007	not measured	0.007	0.006	
DL02-PC05	0.007	not measured	0.007	0.006	a
DL02-PC11	0.528	0.881	0.528	0.478	
DL02-PC12	0.008	not measured	0.008	0.007	a
DL02-PC13	0	3.537	0.000	0.000	
DL02-PC14	13.760	not measured	13.760	12.449	a
DL02-PC15	1.360	0.696	1.360	1.230	c
DL02-PC21	0.005	13.328	0.005	0.004	a
DL02-PC22	0	not measured	0.000	0.000	
DL02-PC23	0.005	not measured	0.005	0.004	a
DL02-PC24	0	not measured	0.000	0.000	

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
DL02-PC25	5.137	not measured	5.137	4.634	a
DL02-PC31	0	0.507	0.000	0.000	
DL02-PC32	0.004	not measured	0.004	0.004	a
DL02-PC33	1.783	not measured	1.783	1.601	
DL02-PC34	0.005	9.214	0.005	0.004	a
DL02-PC35	1.207	not measured	1.207	1.091	c
DL02-PC41	0.006	not measured	0.006	0.005	a
DL02-PC42	0.504	0.001	0.504	0.454	
DL02-PC43	0.007	not measured	0.007	0.007	a
DL02-PC44	0	1.244	0.000	0.000	
DL02-PC45	0.005	0.007	0.005	0.005	a
DL02-PC51	0	not measured	0.000	0.000	
DL02-PC52	0.004	not measured	0.004	0.003	a
DL02-PC53	0	not measured	0.000	0.000	
DL02-PC54	0.004	not measured	0.004	0.004	
DL02-PC55	0	not measured	0.000	0.000	a
DL02-PC61	0.002	not measured	0.002	0.002	a
DL02-PC62	not measured	0.000	0.000	0.000	
GZ01-PC01	0	not measured	0.000	0.000	a
GZ01-PC02	0	not measured	0.000	0.000	
GZ01-PC03	0	not measured	0.000	0.000	a
GZ01-PC04	0	not measured	0.000	0.000	

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
GZ01-PC05	0	not measured	0.000	0.000	a
GZ01-PC11	0	not measured	0.000	0.000	
GZ01-PC12	0	not measured	0.000	0.000	a
GZ01-PC13	0	not measured	0.000	0.000	
GZ01-PC14	not measured	0.042	0.042	0.041	
GZ01-PC15	not measured	0.000	0.000	0.000	
GZ01-PC21	not measured	0.000	0.000	0.000	
GZ01-PC22	not measured	0.000	0.000	0.000	
GZ02-PC01	0	0.000	0.000	0.000	a
GZ02-PC02	0	not measured	0.000	0.000	
GZ02-PC03	0	not measured	0.000	0.000	a
GZ02-PC04	0	not measured	0.000	0.000	
GZ02-PC05	0	not measured	0.000	0.000	a
GZ02-PC11	0	not measured	0.000	0.000	a
GZ02-PC12	57.949	not measured	57.949	56.454	
GZ02-PC13	0	not measured	0.000	0.000	
GZ02-PC14	not measured	43.288	43.288	42.171	
GZ02-PC15	0	not measured	0.000	0.000	
GZ02-PC21	0	not measured	0.000	0.000	a
GZ02-PC22	not measured	0.000	0.000	0.000	
GZ02-PC23	not measured	0.000	0.000	0.000	
GZ02-PC24	not measured	0.000	0.000	0.000	

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
GZ03-PC01	3.664	not measured	3.664	3.560	
GZ03-PC02	32.606	not measured	32.606	31.680	
GZ03-PC03	0	not measured	0.000	0.000	a
GZ03-PC04	0	not measured	0.000	0.000	a
GZ03-PC05	not measured	0.013	0.013	0.013	
GZ03-PC11	not measured	0.024	0.024	0.024	
GZ03-PC12	not measured	0.000	0.000	0.000	
GZ03-PC13	34.132	not measured	34.132	33.163	
GZ03-PC14	6.122	not measured	6.122	5.948	a
GZ03-PC15	0	0.052	0.000	0.000	
GZ03-PC21	0	not measured	0.000	0.000	a
GZ03-PC22	37.673	not measured	37.673	36.603	a
GZ03-PC23	0.024	not measured	0.024	0.024	c
GZ03-PC24	6.647	not measured	6.647	6.471	
GZ03-PC25	not measured	0.000	0.000	0.000	
GZ03-PC31	not measured	0.000	0.000	0.000	
GZ03-PC32	0	not measured	0.000	0.000	a
GZ03-PC33	0.002	not measured	0.002	0.002	a
GZ03-PC34	0	not measured	0.000	0.000	a
GZ03-PC35	0	not measured	0.000	0.000	
GZ03-PC41	0	not measured	0.000	0.000	
GZ03-PC42	not measured	0.000	0.000	0.000	

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
GZ04-PC01	5.504	not measured	5.504	4.999	c
GZ04-PC02	6.009	not measured	6.009	5.458	a,b
GZ04-PC03	32.670	not measured	32.670	31.834	
GZ04-PC04	0	not measured	0.000	0.000	a
GZ04-PC05	not measured	0.142	0.142	0.129	
GZ04-PC11	not measured	0.012	0.012	0.012	
LB01-PC01	40.155	not measured	40.155	32.333	a, c
LB01-PC02	8.774	not measured	8.774	7.065	c
LB02-PC01	0	not measured	0.000	0.000	a
LB02-PC02	0	not measured	0.000	0.000	
LB02-PC03	0.001	not measured	0.001	0.000	a
LB02-PC04	0	not measured	0.000	0.000	
LB03-PC01	49.983	55.301	49.983	39.715	a
LB03-PC02	0.273	not measured	0.273	0.217	
LB04-PC01	26.212	not measured	26.212	20.748	a, c
LB04-PC02	9.820	not measured	9.820	7.773	
LB04-PC03	19.116	not measured	19.116	15.012	a, c
LB04-PC04	9.701	not measured	9.701	7.619	c
LB05-PC01	68.606	not measured	68.606	60.188	a
LB05-PC02	0	not measured	0.000	0.000	
LB05-PC03	78.556	not measured	78.556	68.916	a
LB05-PC04	0	not measured	0.000	0.000	

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
LB06-PC01	3.475	not measured	3.475	2.820	a, c
LB06-PC02	6.630	not measured	6.630	5.380	c
LB06-PC03	7.691	not measured	7.691	6.204	a, c
LB06-PC04	0	not measured	0.000	0.000	
LB06-PC05	22.456	20.773	22.456	18.287	a, c
LB06-PC11	0.288	not measured	0.288	0.235	c
LB07-PC01	36.294	51.347	36.294	28.685	a, c
LB07-PC02	64.752	not measured	64.752	51.178	
LB07-PC03	3.293	not measured	3.293	2.612	a, c
LB07-PC04	26.980	33.879	26.980	21.399	
OF01-PC01	7.125	not measured	7.125	6.864	
OF02-PC01	0	not measured	0.000	0.000	
OF03-PC01	1.556	not measured	1.556	1.499	c
OF04-PC01	0	not measured	0.000	0.000	
OF05-PC01	1.607	not measured	1.607	1.548	c
OF06-PC01	0	not measured	0.000	0.000	
OF07-PC01	6.615	not measured	6.615	6.373	
OF08-PC01	0	not measured	0.000	0.000	
OF08-PC02	0	not measured	0.000	0.000	
OF09-PC01	0	not measured	0.000	0.000	
OF10-PC01	2.696	not measured	2.696	2.597	
OF11-PC01	6.589	not measured	6.589	6.347	

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
OF11-PC02	2.821	not measured	2.821	2.717	
OF11-PC03	0	not measured	0.000	0.000	
OF11-PC04	0	not measured	0.000	0.000	
RB01-PC01	0	not measured	0.000	0.000	
RB01-PC02	0	not measured	0.000	0.000	
RB01-PC03	0	not measured	0.000	0.000	
RB01-PC04	7.787	not measured	7.787	6.400	
RB01-PC05	0.123	not measured	0.123	0.101	
RB01-PC11	0.199	not measured	0.199	0.163	
RB01-PC12	0	not measured	0.000	0.000	
RB01-PC13	17.226	not measured	17.226	14.159	
RB01-PC14	4.163	0.194	4.163	3.422	
RB01-PC15	0	not measured	0.000	0.000	
RB01-PC21	15.169	not measured	15.169	12.468	
RB01-PC23	19.824	not measured	19.824	16.294	
RB01-PC24	0	not measured	0.000	0.000	
RB01-PC25	0	not measured	0.000	0.000	
RB01-PC34	16.795	not measured	16.795	13.804	
RB02-PC01	0	6.664	0.000	0.000	
RB02-PC02	0	not measured	0.000	0.000	
RB02-PC03	9.578	not measured	9.578	6.930	
RB02-PC05	7.949	not measured	7.949	5.751	c

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
RB02-PC11	0.002	not measured	0.002	0.001	
RB02-PC12	0.004	not measured	0.004	0.003	
RQ01-PC01	0	not measured	0.000	0.000	a
RQ02-PC01	6.316	not measured	6.316	5.446	a
RQ02-PC02	0.001	not measured	0.001	0.001	
RQ03-PC01	19.088	NA	19.088	16.341	a
RQ04-PC01	0	not measured	0.000	0.000	a
RQ05-PC01	0	not measured	0.000	0.000	
RQ05-PC02	0.008	0.515	0.008	0.007	a
RQ05-PC03	17.390	not measured	17.390	15.253	
RQ05-PC04	0	not measured	0.000	0.000	a
RQ06-PC01	0	not measured	0.000	0.000	a
RQ06-PC02	0	not measured	0.000	0.000	
RQ06-PC03	0	0.000	0.000	0.000	a
RQ06-PC04	0	not measured	0.000	0.000	
RQ07-PC01	0	not measured	0.000	0.000	
RQ07-PC02	0	not measured	0.000	0.000	
RQ07-PC03	18.241	20.992	18.241	14.179	a
RQ07-PC04	9.078	not measured	9.078	7.056	a
RQ07-PC05	0	not measured	0.000	0.000	a
VF01-PC01	2.347	not measured	2.347	2.288	
VF01-PC02	0.003	not measured	0.003	0.003	a

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
VF01-PC03	0	not measured	0.000	0.000	
VF01-PC04	0	not measured	0.000	0.000	a
VF01-PC05	5.387	not measured	5.387	5.259	a
VF01-PC11	0	not measured	0.000	0.000	
VF01-PC12	0.001	not measured	0.001	0.001	a
VF01-PC13	0	not measured	0.000	0.000	
VF01-PC14	0.001	not measured	0.001	0.001	a
VF01-PC15	0	not measured	0.000	0.000	
VF01-PC21	0	not measured	0.000	0.000	a
VF01-PC22	16.807	not measured	16.807	16.405	
VF01-PC23	0	not measured	0.000	0.000	a
VF01-PC24	0	not measured	0.000	0.000	
VF01-PC25	0	not measured	0.000	0.000	
VF01-PC31	0.001	not measured	0.001	0.001	a
VF01-PC32	0.001	not measured	0.001	0.001	a
VF01-PC33	0	not measured	0.000	0.000	
VF01-PC34	0	not measured	0.000	0.000	
VF01-PC35	0.001	not measured	0.001	0.001	a
VF01-PC41	0	not measured	0.000	0.000	a
VF01-PC42	0	not measured	0.000	0.000	
VF01-PC43	0	not measured	0.000	0.000	a
VF01-PC44	0	not measured	0.000	0.000	

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
VF01-PC45	8.505	not measured	8.505	8.291	
VF01-PC51	11.023	not measured	11.023	10.761	a
VF01-PC52	0	not measured	0.000	0.000	a
VF01-PC53	9.645	not measured	9.645	9.427	
VF01-PC54	4.544	not measured	4.544	4.436	
VF01-PC55	0.002	not measured	0.002	0.002	a
VF02-PC01	0	not measured	0.000	0.000	a
VF02-PC02	0	not measured	0.000	0.000	
VF02-PC03	1.119	not measured	1.119	1.083	a
VF02-PC04	0	not measured	0.000	0.000	
VF02-PC05	0	not measured	0.000	0.000	a
VF02-PC11	0	not measured	0.000	0.000	
VF02-PC12	0	not measured	0.000	0.000	
XQ01-PC01	0	not measured	0.000	0.000	a
XQ01-PC02	0	not measured	0.000	0.000	
XQ01-PC03	not measured	0.000	0.000	0.000	
XQ01-PC04	37.374	0.125	37.374	32.969	a
XQ02-PC01	0	not measured	0.000	0.000	a
XQ02-PC02	0	0.000	0.000	0.000	
XQ02-PC03	not measured	0.000	0.000	0.000	
XQ02-PC04	14.982	not measured	14.982	12.952	a
XQ03-PC01	14.955	not measured	14.955	13.587	a

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
XQ03-PC02	16.531	not measured	16.531	15.019	
XQ03-PC03	not measured	0.000	0.000	0.000	
XQ03-PC04	14.500	not measured	14.500	13.174	a
XQ04-PC01	0	not measured	0.000	0.000	a
XQ04-PC02	37.406	not measured	37.406	33.473	
XQ04-PC03	not measured	0.000	0.000	0.000	
XQ04-PC04	not measured	2.866	2.866	2.564	
XQ05-PC01	0	not measured	0.000	0.000	a
XQ05-PC02	24.085	not measured	24.085	21.714	
XQ05-PC03	not measured	0.000	0.000	0.000	
XQ05-PC04	not measured	0.000	0.000	0.000	
XQ06-PC01	20.160	not measured	20.160	17.970	a
XQ06-PC02	11.918	not measured	11.918	10.623	
XQ06-PC03	not measured	0.000	0.000	0.000	
XQ06-PC04	not measured	0.149	0.149	0.133	
XQ07-PC01	0	not measured	0.000	0.000	a
XQ07-PC02	0	not measured	0.000	0.000	
XQ07-PC03	not measured	0.000	0.000	0.000	
XQ07-PC04	not measured	1.752	1.752	1.605	
ZW01-PC01	6.149	not measured	6.149	5.572	
ZW01-PC02	0	not measured	0.000	0.000	
ZW01-PC03	0	not measured	0.000	0.000	

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
ZW01-PC04	0	not measured	0.000	0.000	
ZW01-PC05	0	not measured	0.000	0.000	
ZW01-PC11	0	not measured	0.000	0.000	
ZW01-PC12	0	not measured	0.000	0.000	
ZW01-PC13	2.664	not measured	2.664	2.421	
ZW01-PC14	0	not measured	0.000	0.000	
ZW01-PC15	0	not measured	0.000	0.000	
ZW01-PC21	3.058	not measured	3.058	2.779	c
ZW01-PC22	0	not measured	0.000	0.000	
ZW01-PC23	0	not measured	0.000	0.000	
ZW01-PC24	0	not measured	0.000	0.000	
ZW01-PC25	0	not measured	0.000	0.000	
ZW01-PC31	0	not measured	0.000	0.000	
ZW01-PC32	0	not measured	0.000	0.000	
ZW01-PC33	0	not measured	0.000	0.000	
ZW01-PC34	0	not measured	0.000	0.000	
ZW01-PC35	0	not measured	0.000	0.000	
ZW02-PC01	11.284	not measured	11.284	10.116	
ZW02-PC02	0	not measured	0.000	0.000	
ZW02-PC03	0	not measured	0.000	0.000	
ZW02-PC04	0	not measured	0.000	0.000	
ZW02-PC12	0	not measured	0.000	0.000	

Table A-8 (continued).

Device Identifier	Supply Gas meter - avg. emission rate whole gas (scf/h)	Hi Flow® - avg. emission rate whole gas (scf/h)	Whole gas flow rate used in data analysis (scf/h)	Methane flow rate used in data analysis (scf/h)	Flags
ZW03-PC01	0	not measured	0.000	0.000	
ZW03-PC02	3.590	not measured	3.590	3.275	
ZW03-PC03	0	not measured	0	0	

Flags: (a): Correction factor was applied to account for the condensation of an oily substance on the sensor of the measuring instrument.

(b): Rapid peak in time series attributed to voltage surge associated with vehicle, which was the source of power for the supply gas meters used in the study, being turned on/off rather than with a pneumatic controller actuation.

(c): Devices used in volume per actuation analysis for the “zero” pattern emissions.

Table A-9. Total measurement time, number of actuations, and frequency of actuations.

Device Identifier	Total measured time (min)	# of actuations	frequency of actuations (#/min)
AA01-PC04	15.4	51	3.3
AA01-PC05	109.2	0	no actuations
AA01-PC06	23.0	24	1.0
AA01-PC07	18.7	1	0.1
AA01-PC08	15.2	10	0.7
AA01-PC09	15.7	2	0.1
AA01-PC11	14.3	2	0.1
AA02-PC04	11.0	0	no actuations
AA02-PC05	14.8	2	0.1
AA02-PC06	15.7	1	0.1
AA02-PC07	18.0	4	0.2
AA02-PC08	10.3	0	no actuations
AA02-PC09	11.2	0	no actuations
AP01-PC01	15.7	12	0.8
AP01-PC02	14.3	0	no actuations
AP01-PC03	53.1	0	no actuations
AP01-PC04	15.0	0	no actuations
AP01-PC05	15.0	0	no actuations
AP01-PC12	15.0	78	5.2
AP02-PC01	15.1	4	0.3
AP02-PC02	15.4	10	0.6
AP02-PC03	16.0	0	no actuations
AP02-PC04	15.0	7	0.5
AP03-PC01	16.7	0	no actuations
AP03-PC02	16.1	0	no actuations
AP04-PC01	16.0	0	no actuations
AP04-PC02	16.6	35	2.1
AP04-PC03	13.5	5	0.4
AP04-PC04	15.1	0	no actuations
AP05-PC01	23.9	6	0.3
AP05-PC02	14.0	0	no actuations
CW01-PC01	15.0	0	no actuations
CW01-PC02	15.0	0	no actuations
CW01-PC03	15.7	0	no actuations
CW01-PC04	15.5	0	no actuations
CW01-PC05	15.6	0	no actuations
CW01-PC11	19.5	2	0.1

Table A-9 (continued).

Device Identifier	Total measured time (min)	# of actuations	frequency of actuations (#/min)
CW01-PC12	14.8	1	0.1
CW01-PC13	15.5	1	0.1
CW01-PC14	15.7	1	0.1
CW01-PC15	15.0	0	no actuations
CW01-PC21	14.7	0	no actuations
CW01-PC22	15.7	0	no actuations
CW01-PC23	15.7	21	1.3
CW01-PC24	15.8	0	no actuations
CW01-PC25	15.6	0	no actuations
CW01-PC31	15.3	0	no actuations
CW01-PC32	18.5	0	no actuations
CW01-PC33	15.3	0	no actuations
CW01-PC34	15.4	0	no actuations
CW01-PC35	18.5	0	no actuations
CW02-PC01	15.0	0	no actuations
CW02-PC02	15.3	0	no actuations
CW02-PC03	14.6	0	no actuations
CW02-PC04	14.5	0	no actuations
CW02-PC05	15.4	0	no actuations
CW02-PC11	1.50	0	no actuations
CW02-PC12	15.4	0	no actuations
CW02-PC13	16.8	0	no actuations
CW02-PC14	16.9	0	no actuations
CW02-PC15	15.5	0	no actuations
CW02-PC21	1.50	0	no actuations
CW02-PC22	15.3	0	no actuations
CW02-PC23	15.3	0	no actuations
CW02-PC24	15.2	0	no actuations
CW02-PC25	15.2	0	no actuations
CW02-PC31	15.0	0	no actuations
CW02-PC32	16.3	3	0.2
CW02-PC33	20.2	1	0.0
CW02-PC34	20.2	0	no actuations
CW02-PC35	16.3	0	no actuations
CZ01-PC01	14.9	1	0.1
CZ01-PC02	15.0	0	no actuations
CZ02-PC01	15.2	0	no actuations

Table A-9 (continued).

Device Identifier	Total measured time (min)	# of actuations	frequency of actuations (#/min)
CZ02-PC02	15.0	1	0.7
CZ03-PC01	16.4	0	no actuations
CZ03-PC02	15.0	0	no actuations
CZ04-PC01	16.1	0	no actuations
CZ04-PC02	15.0	0	no actuations
CZ05-PC01	15.3	0	no actuations
CZ05-PC02	15.0	0	no actuations
CZ05-PC03	15.3	0	no actuations
CZ06-PC01	15.0	0	no actuations
CZ06-PC02	15.0	0	no actuations
CZ06-PC03	15.0	0	no actuations
CZ07-PC01	16.2	0	no actuations
CZ07-PC02	15.0	0	no actuations
CZ08-PC01	15.1	0	no actuations
CZ08-PC02	16.1	3	0.2
CZ09-PC01	15.1	0	no actuations
CZ09-PC02	15.2	0	no actuations
CZ10-PC01	29.5	2	0.1
CZ11-PC01	15.7	0	no actuations
DL01-PC01	15.0	1	0.1
DL01-PC02	15.0	0	no actuations
DL01-PC03	14.9	0	no actuations
DL01-PC04	15.0	0	no actuations
DL01-PC05	14.4	0	no actuations
DL01-PC11	14.8	0	no actuations
DL01-PC12	14.7	1	0.1
DL01-PC13	14.9	0	no actuations
DL01-PC14	14.8	1	0.1
DL01-PC15	14.9	1	0.1
DL01-PC21	14.8	7	0.5
DL01-PC22	14.7	0	no actuations
DL01-PC23	14.7	7	0.5
DL01-PC24	15.5	0	no actuations
DL01-PC25	15.5	0	no actuations
DL01-PC31	14.5	0	no actuations
DL01-PC32	15.0	0	no actuations
DL01-PC33	15.0	1	0.1

Table A-9 (continued).

Device Identifier	Total measured time (min)	# of actuations	frequency of actuations (#/min)
DL01-PC34	15.2	0	no actuations
DL01-PC35	15.2	0	no actuations
DL01-PC41	15.7	0	no actuations
DL01-PC42	15.7	1	0.1
DL01-PC43	14.9	0	no actuations
DL01-PC44	14.9	1	0.1
DL01-PC45	15.0	0	no actuations
DL02-PC01	14.9	0	no actuations
DL02-PC02	14.9	1	0.1
DL02-PC03	14.9	0	no actuations
DL02-PC04	14.9	1	0.1
DL02-PC05	14.9	0	no actuations
DL02-PC11	14.8	1	0.1
DL02-PC12	15.2	0	no actuations
DL02-PC13	15.3	0	no actuations
DL02-PC14	14.8	30	2.0
DL02-PC15	14.9	30	2.0
DL02-PC21	16.1	0	no actuations
DL02-PC22	16.0	0	no actuations
DL02-PC23	15.0	0	no actuations
DL02-PC24	14.7	0	no actuations
DL02-PC25	15.3	0	no actuations
DL02-PC31	15.2	0	no actuations
DL02-PC32	14.9	0	no actuations
DL02-PC33	14.8	33	2.2
DL02-PC34	15.2	0	no actuations
DL02-PC35	15.1	11	0.7
DL02-PC41	14.7	0	no actuations
DL02-PC42	14.7	1	0.1
DL02-PC43	15.0	0	no actuations
DL02-PC44	15.0	0	no actuations
DL02-PC45	14.9	0	no actuations
DL02-PC51	14.9	0	no actuations
DL02-PC52	15.1	0	no actuations
DL02-PC53	15.1	0	no actuations
DL02-PC54	14.6	0	no actuations
DL02-PC55	14.6	0	no actuations

Table A-9 (continued).

Device Identifier	Total measured time (min)	# of actuations	frequency of actuations (#/min)
DL02-PC61	14.5	0	no actuations
DL02-PC62	15.0	0	no actuations
GZ01-PC01	15.3	0	no actuations
GZ01-PC02	15.7	0	no actuations
GZ01-PC03	15.2	0	no actuations
GZ01-PC04	15.1	0	no actuations
GZ01-PC05	15.4	0	no actuations
GZ01-PC11	15.3	0	no actuations
GZ01-PC12	14.6	0	no actuations
GZ01-PC13	14.4	0	no actuations
GZ01-PC14	15.0	0	no actuations
GZ01-PC15	15.0	0	no actuations
GZ01-PC21	15.0	0	no actuations
GZ01-PC22	15.0	0	no actuations
GZ02-PC01	14.6	0	no actuations
GZ02-PC02	14.5	0	no actuations
GZ02-PC03	14.8	0	no actuations
GZ02-PC04	15.6	0	no actuations
GZ02-PC05	15.0	0	no actuations
GZ02-PC11	15.2	0	no actuations
GZ02-PC12	15.2	0	no actuations
GZ02-PC13	15.0	0	no actuations
GZ02-PC14	15.0	0	no actuations
GZ02-PC15	15.1	0	no actuations
GZ02-PC21	15.3	0	no actuations
GZ02-PC22	15.0	0	no actuations
GZ02-PC23	15.0	0	no actuations
GZ02-PC24	15.0	0	no actuations
GZ03-PC01	15.2	2	0.1
GZ03-PC02	15.5	2	0.1
GZ03-PC03	15.5	0	no actuations
GZ03-PC04	15.2	0	no actuations
GZ03-PC05	15.0	0	no actuations
GZ03-PC11	15.0	0	no actuations
GZ03-PC12	15.0	0	no actuations
GZ03-PC13	17.6	0	no actuations
GZ03-PC14	17.6	0	no actuations

Table A-9 (continued).

Device Identifier	Total measured time (min)	# of actuations	frequency of actuations (#/min)
GZ03-PC15	14.9	0	no actuations
GZ03-PC21	90.5	0	no actuations
GZ03-PC22	15.0	0	no actuations
GZ03-PC23	90.4	14	0.2
GZ03-PC24	15.2	0	no actuations
GZ03-PC25	15.0	0	no actuations
GZ03-PC31	15.0	0	no actuations
GZ03-PC32	14.9	0	no actuations
GZ03-PC33	14.9	0	no actuations
GZ03-PC34	14.9	0	no actuations
GZ03-PC35	14.9	0	no actuations
GZ03-PC41	15.1	0	no actuations
GZ03-PC42	15.0	0	no actuations
GZ04-PC01	16.0	2	0.1
GZ04-PC02	16.0	0	no actuations
GZ04-PC03	17.8	5	0.3
GZ04-PC04	17.9	0	no actuations
GZ04-PC05	15.0	0	no actuations
GZ04-PC11	15.0	0	no actuations
LB01-PC01	15.0	4	0.3
LB01-PC02	15.0	3	0.2
LB02-PC01	15.0	0	no actuations
LB02-PC02	15.0	0	no actuations
LB02-PC03	15.0	0	no actuations
LB02-PC04	15.0	0	no actuations
LB03-PC01	15.2	8	0.5
LB03-PC02	15.2	1	0.1
LB04-PC01	15.0	5	0.3
LB04-PC02	15.0	6	0.4
LB04-PC03	20.2	8	0.4
LB04-PC04	20.2	2	0.1
LB05-PC01	15.1	11	0.7
LB05-PC02	15.1	0	no actuations
LB05-PC03	15.5	15	1.0
LB05-PC04	15.5	0	no actuations
LB06-PC01	15.0	4	0.3
LB06-PC02	15.0	2	0.1

Table A-9 (continued).

Device Identifier	Total measured time (min)	# of actuations	frequency of actuations (#/min)
LB06-PC03	15.2	11	0.7
LB06-PC04	15.2	0	no actuations
LB06-PC05	15.7	14	0.9
LB06-PC11	15.7	6	0.4
LB07-PC01	15.0	15	1.0
LB07-PC02	15.0	15	1.0
LB07-PC03	18.1	14	0.8
LB07-PC04	21.1	2	0.1
OF01-PC01	15.0	23	1.5
OF02-PC01	14.9	0	no actuations
OF03-PC01	15.0	154	10.3
OF04-PC01	15.0	0	no actuations
OF05-PC01	15.4	6	0.4
OF06-PC01	15.0	0	no actuations
OF07-PC01	15.0	98	6.5
OF08-PC01	15.2	0	no actuations
OF08-PC02	15.0	0	no actuations
OF09-PC01	15.1	0	no actuations
OF10-PC01	15.1	0	no actuations
OF11-PC01	22.7	10	0.4
OF11-PC02	14.9	0	no actuations
OF11-PC03	15.7	0	no actuations
OF11-PC04	15.3	0	no actuations
RB01-PC01	15.0	0	no actuations
RB01-PC02	11.0	0	no actuations
RB01-PC03	15.0	0	no actuations
RB01-PC04	14.9	0	no actuations
RB01-PC05	18.0	0	no actuations
RB01-PC11	18.0	1	0.1
RB01-PC12	16.0	0	no actuations
RB01-PC13	14.9	0	no actuations
RB01-PC14	16.9	8	0.5
RB01-PC15	15.0	0	no actuations
RB01-PC21	15.0	0	no actuations
RB01-PC23	15.0	0	no actuations
RB01-PC24	15.0	0	no actuations
RB01-PC25	15.0	0	no actuations

Table A-9 (continued).

Device Identifier	Total measured time (min)	# of actuations	frequency of actuations (#/min)
RB01-PC34	15.0	0	no actuations
RB02-PC01	15.0	0	no actuations
RB02-PC02	15.8	0	no actuations
RB02-PC03	15.0	0	no actuations
RB02-PC05	15.0	5	0.3
RB02-PC11	15.0	0	no actuations
RB02-PC12	15.0	0	no actuations
RQ01-PC01	15.1	0	no actuations
RQ02-PC01	16.2	0	no actuations
RQ02-PC02	16.1	0	no actuations
RQ03-PC01	20.4	0	no actuations
RQ04-PC01	39.9	0	no actuations
RQ05-PC01	20.2	0	no actuations
RQ05-PC02	20.3	1	0.0
RQ05-PC03	14.8	0	no actuations
RQ05-PC04	15.0	0	no actuations
RQ06-PC01	14.8	0	no actuations
RQ06-PC02	15.0	0	no actuations
RQ06-PC03	15.9	0	no actuations
RQ06-PC04	15.8	0	no actuations
RQ07-PC01	15.1	0	no actuations
RQ07-PC02	15.0	0	no actuations
RQ07-PC03	16.1	0	no actuations
RQ07-PC04	15.2	0	no actuations
RQ07-PC05	15.2	0	no actuations
VF01-PC01	15.7	0	no actuations
VF01-PC02	15.8	0	no actuations
VF01-PC03	15.3	0	no actuations
VF01-PC04	15.4	0	no actuations
VF01-PC05	15.2	0	no actuations
VF01-PC11	15.2	0	no actuations
VF01-PC12	15.8	0	no actuations
VF01-PC13	15.9	0	no actuations
VF01-PC14	14.1	0	no actuations
VF01-PC15	14.1	0	no actuations
VF01-PC21	15.2	0	no actuations
VF01-PC22	15.2	0	no actuations

Table A-9 (continued).

Device Identifier	Total measured time (min)	# of actuations	frequency of actuations (#/min)
VF01-PC23	15.2	0	no actuations
VF01-PC24	15.2	0	no actuations
VF01-PC25	15.2	0	no actuations
VF01-PC31	15.2	0	no actuations
VF01-PC32	15.0	0	no actuations
VF01-PC33	16.1	0	no actuations
VF01-PC34	15.1	0	no actuations
VF01-PC35	15.1	0	no actuations
VF01-PC41	17.1	0	no actuations
VF01-PC42	17.1	0	no actuations
VF01-PC43	14.9	0	no actuations
VF01-PC44	14.9	0	no actuations
VF01-PC45	14.5	0	no actuations
VF01-PC51	14.5	0	no actuations
VF01-PC52	15.2	0	no actuations
VF01-PC53	15.2	1	0.1
VF01-PC54	16.0	1	0.1
VF01-PC55	16.0	0	no actuations
VF02-PC01	15.3	0	no actuations
VF02-PC02	15.3	0	no actuations
VF02-PC03	15.0	1	0.1
VF02-PC04	15.0	0	no actuations
VF02-PC05	14.6	0	no actuations
VF02-PC11	14.6	0	no actuations
VF02-PC12	15.0	0	no actuations
XQ01-PC01	15.0	0	no actuations
XQ01-PC02	15.0	0	no actuations
XQ01-PC03	15.0	0	no actuations
XQ01-PC04	15.3	3	0.2
XQ02-PC01	15.9	0	no actuations
XQ02-PC02	15.9	0	no actuations
XQ02-PC03	15.0	0	no actuations
XQ02-PC04	14.0	0	no actuations
XQ03-PC01	15.0	0	no actuations
XQ03-PC02	15.0	0	no actuations
XQ03-PC03	15.0	0	no actuations
XQ03-PC04	15.0	0	no actuations

Table A-9 (continued).

Device Identifier	Total measured time (min)	# of actuations	frequency of actuations (#/min)
XQ04-PC01	15.9	0	no actuations
XQ04-PC02	15.9	0	no actuations
XQ04-PC03	15.0	0	no actuations
XQ04-PC04	15.0	0	no actuations
XQ05-PC01	15.0	0	no actuations
XQ05-PC02	15.0	0	no actuations
XQ05-PC03	15.0	0	no actuations
XQ05-PC04	15.0	0	no actuations
XQ06-PC01	15.0	0	no actuations
XQ06-PC02	15.0	0	no actuations
XQ06-PC03	15.0	0	no actuations
XQ06-PC04	15.0	0	no actuations
XQ07-PC01	17.0	0	no actuations
XQ07-PC02	17.0	0	no actuations
XQ07-PC03	15.0	0	no actuations
XQ07-PC04	15.0	1	0.7
ZW01-PC01	15.3	0	no actuations
ZW01-PC02	15.0	0	no actuations
ZW01-PC03	15.0	0	no actuations
ZW01-PC04	15.3	0	no actuations
ZW01-PC05	15.4	0	no actuations
ZW01-PC11	14.9	0	no actuations
ZW01-PC12	22.0	0	no actuations
ZW01-PC13	15.0	0	no actuations
ZW01-PC14	15.5	0	no actuations
ZW01-PC15	14.9	0	no actuations
ZW01-PC21	15.0	4	0.3
ZW01-PC22	14.8	0	no actuations
ZW01-PC23	14.9	0	no actuations
ZW01-PC24	15.0	0	no actuations
ZW01-PC25	14.9	0	no actuations
ZW01-PC31	15.0	0	no actuations
ZW01-PC32	15.0	0	no actuations
ZW01-PC33	14.9	0	no actuations
ZW01-PC34	14.9	0	no actuations
ZW01-PC35	15.1	0	no actuations
ZW02-PC01	15.0	0	no actuations

Table A-9 (continued).

Device Identifier	Total measured time (min)	# of actuations	frequency of actuations (#/min)
ZW02-PC02	14.9	0	no actuations
ZW02-PC03	15.0	0	no actuations
ZW02-PC04	15.0	0	no actuations
ZW02-PC12	15.0	0	no actuations
ZW03-PC01	15.0	0	no actuations
ZW03-PC02	15.0	0	no actuations
ZW03-PC03	15.0	0	no actuations

A.5 Estimates of emissions from devices with no emissions detected during 15 minute sampling period

The data set includes 241 devices (64%) for which no emissions were detected over a 15 minute sampling period (136 devices) or that had flow below the instrument measurement threshold that was indistinguishable from instrument noise (105 devices). Even though no actuations were detected on these devices during the sampling period, for some of these devices, actuations and the associated emissions per actuation would have been observed if the sampling period had been extended. Estimates of the potential emissions from these devices can be determined by estimating actuation frequency and volume.

$$\text{Estimated emissions} = \text{actuation frequency (min}^{-1}\text{)} * \text{actuation volume (scf)} \quad (\text{A5.1})$$

It is assumed, in these estimates, that all of these devices for which emissions are estimated are intermittent vent controllers, rather than continuous vent controllers, since the measured emission rates included zero values.

The actuation frequencies and emissions per actuation were estimated based on data for controllers with well-defined actuation patterns. A well-defined actuation pattern exhibited intermittent emissions patterns, returned to a zero emissions baseline between actuations, and had two or more actuations during the 15 minute sampling period. Average actuation volumes for controllers in eight types of service are provided in Table A-10.

Table A-10. Counts of devices and average emissions per actuation for controllers in eight application categories; note that for ESDs, the emissions measured during the sampling were device leak rates rather than device actuations.

Application	Devices with no emissions detected during sampling period	Devices with well-defined actuations	Average volume per actuation for “Actuating” Devices (scf)
Separator - Level Control	109	26	0.349
ESD	31	5	0.026
Plunger Lift	20	1	0.048
Process Heater	40	1	0.191
Other	10	0	Assumed average from all other devices (0.271)
Separator - Other	9	2	0.070
Compressor	7	2	0.158
Dehydration System	15	1	0.398

Table A-10 provides estimates of actuation volumes, categorized by application. Actuation frequency could be estimated using a variety of approaches. An upper bound on the emission rate could be estimated by assuming that all devices for which no emissions were observed over 15 minutes of sampling actually had actuations that occurred immediately before and immediately after the sampling period (4 actuations per hour, a frequency of 0.067 min^{-1}). This scenario is improbable, nevertheless, it does provide an upper bound. Assuming that devices with no measured emissions actuated every 15 minutes, and had the actuation volumes, consistent with application type, reported in Table A-10, increases the overall population average emissions by 11% (from 5.52 scf/h to 6.13 scf/h whole gas). As the assumed actuation frequency decreases, the estimated additional emissions decrease, as shown in Figure A-5. If a 3 hour period between actuations is assumed (0.0055 min^{-1}), the study average emissions rate per device increases by less than 1%.

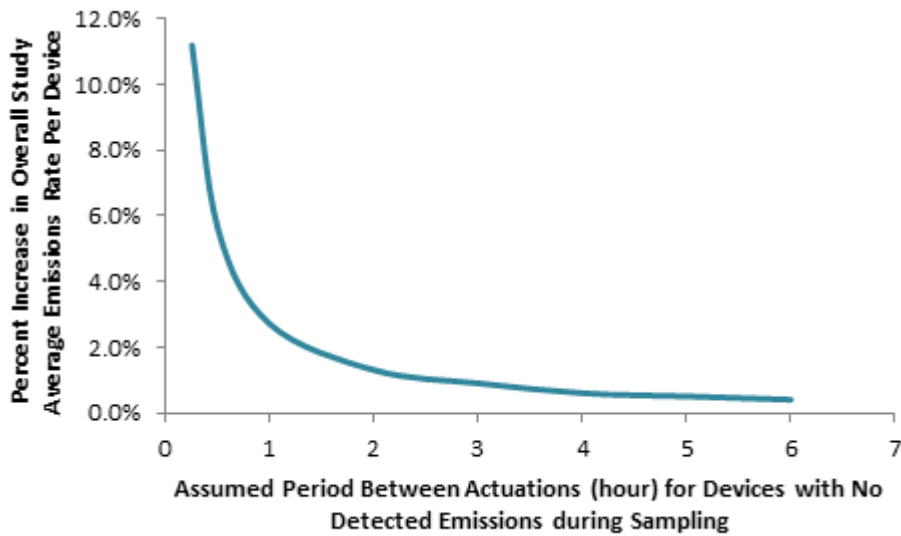


Figure A-5. Sensitivity of study-average emissions rate per device to the assumed period between actuations for devices with no detected emissions during the sampling period.

For some application categories, specifically level control of separators, actuation frequencies, for devices with no actuations detected over 15 minutes of sampling, can be estimated based on the data from devices with observed actuations. Separator level controllers were both the largest category of pneumatic devices measured in this study (50% of total devices) and the largest category of devices with no measured emissions over a 15 minute sampling period (45%). For this category of devices, the actuation frequencies for all intermittent vent controllers with a period between actuations of less than 7.5 minutes (at least two actuations in 15 minutes of observation, 21% of the intermittent vent separator level controllers) are shown in Figure A-6. The most conservative estimate of the range of periods between actuations for separator level controllers would be a linear extrapolation of the subset of devices with a periodicity of less than 7.5 minutes (Figure A-6).

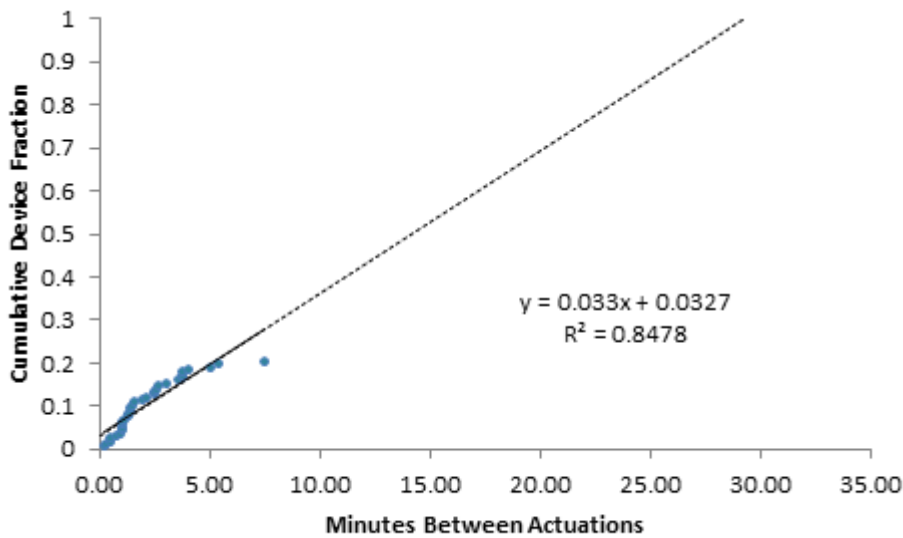


Figure A-6. Period between actuations for separator level controllers with intermittent vent emissions patterns.

A linear extrapolation of the actuation frequency for separator level controllers with two or more actuations in the sample period to the overall population of separator level controllers indicated that all separator level controllers would undergo an actuation within 30 minutes. Assuming that the minutes between actuations for the separator level controllers for which no emissions were detected during sampling were evenly spaced between 15 minutes and 30 minutes, the increase in the overall study-average per device emissions rate for all devices in all applications would be between 5.1% (assuming that the actual emissions rate from all non-separator level controllers for which no emissions were detected during sampling were negligible) and 8.9% (assuming that all devices without detected emissions during sampling and that were not in separator level controller applications actuated every 15 minutes). This approach likely under-estimates the time between actuations for separator level controllers (over-estimates the frequency). Based on the linear extrapolation shown in Figure A-6-, 25% of devices in separator level control applications would be expected to have actuations between 7.5 minutes to 15 minutes. For devices sampled in the study, only 6% of level controllers on separators recorded a single actuation during a 15 minute sample time or had a periodicity in this

range due to two actuations during a sample time longer than 15 minutes. Adjusting the slope of the linear interpolation to fit the data collected between 7.5 and 15 minutes changes the slope to 0.008 and would indicate that all devices would actuate within 121 minutes. Separator level controller emissions would be estimated to increase by 2.9% under this assumption. Under this revised linear interpolation, the estimated average emissions per device from the overall study population could increase between 2.1% and 5.9%, again depending on the frequency assumed for controllers not in separator level control service.

Overall, the study average emission rate for controllers would be expected to increase by 2%-6% if the measurement period had been extended indefinitely. This estimate is based on data for controllers in separator level control service, the most common type of service observed in the study, and a type of service that is likely to result in regular actuations.

A.6 Numbers of controllers per well

In this work, a total of 65 sites were visited. The 65 sites had 377 pneumatic controllers on which measurements were performed and 53 for which either time constraints or equipment failure led to no sample being collected. The 65 sites with 430 controllers had 161 wells, for an average of 2.7 devices per well. The median value was 2.0 controllers per well with 25th and 75th percentile values of 1 and 4 controllers per well. The minimum value was 0.4 devices per well, and the maximum value was 11 devices per well. The distribution is shown in Figure A-7. Table A-11 provides counts of controllers and wells at individual sites.

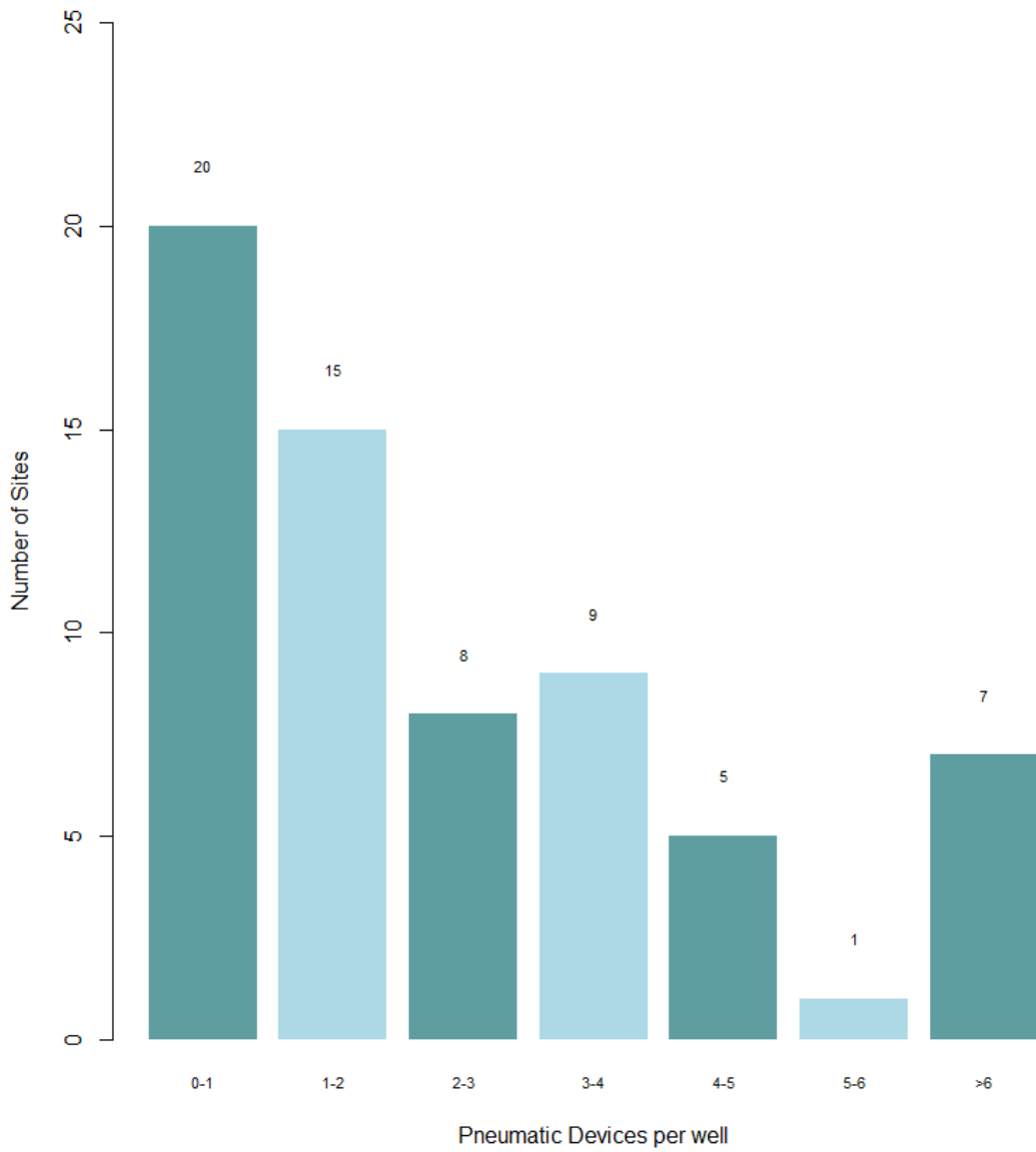


Figure A-7. Distribution of number of pneumatic devices per well for production sites sampled by the study team.

Table A-11. Number of pneumatic devices and wells for sites sampled in this work.

Site Identifier	Count of Measured Pneumatic Devices	Count of Not measured Pneumatic Devices:	Total Pneumatic Devices	Count of wells	Devices/well
AA01	7	22	29	3	9.7
AA02	6	14	20	2	10.0
AP01	6	0	6	16	0.4
AP02	4	0	4	10	0.4
AP03	2	0	2	3	0.7
AP04	4	0	4	8	0.5
AP05	2	0	2	4	0.5
CW01	20	0	20	4	5.0
CW02	20	0	20	4	5.0
CZ01	2	0	2	1	2.0
CZ02	2	0	2	1	2.0
CZ03	2	0	2	1	2.0
CZ04	2	0	2	1	2.0
CZ05	3	0	3	1	3.0
CZ06	3	0	3	1	3.0
CZ07	2	0	2	1	2.0
CZ08	2	1	3	3	1.0
CZ09	2	0	2	1	2.0
CZ10	1	0	1	1	1.0
CZ11	1	0	1	1	1.0
DL01	25	0	25	10	2.5
DL02	32	0	32	14	2.3
GZ01	12	0	12	4	3.0
GZ02	14	0	14	2	7.0
GZ03	22	0	22	2	11.0
GZ04	6	0	6	2	3.0
LB01	2	0	2	1	2.0
LB02	4	0	4	2	2.0
LB03	2	0	2	1	2.0
LB04	4	0	4	2	2.0
LB05	4	2	6	2	3.0
LB06	6	3	9	3	3.0
LB07	4	0	4	2	2.0
OF01	1	0	1	1	1.0
OF02	1	0	1	1	1.0
OF03	1	0	1	1	1.0

Table A-11 (continued).

Site Identifier	Count of Measured Pneumatic Devices	Count of Not measured Pneumatic Devices:	Total Pneumatic Devices	Count of wells	Devices/well
OF04	1	0	1	1	1.0
OF05	1	0	1	1	1.0
OF06	1	0	1	1	1.0
OF07	1	0	1	1	1.0
OF08	2	0	2	1	2.0
OF09	1	0	1	1	1.0
OF10	1	0	1	1	1.0
OF11	4	0	4	1	4.0
RB01	15	0	15	3	5.0
RB02	6	1	7	1	7.0
RQ01	1	0	1	1	1.0
RQ02	2	0	2	1	2.0
RQ03	1	0	1	1	1.0
RQ04	1	0	1	1	1.0
RQ05	4	0	4	2	2.0
RQ06	4	0	4	1	4.0
RQ07	5	0	5	1	5.0
VF01	30	0	30	6	5.0
VF02	7	0	7	6	1.2
XQ01	4	0	4	1	4.0
XQ02	4	0	4	1	4.0
XQ03	4	0	4	1	4.0
XQ04	4	0	4	1	4.0
XQ05	4	0	4	1	4.0
XQ06	4	0	4	1	4.0
XQ07	4	0	4	1	4.0
ZW01	20	5	25	3	8.3
ZW02	5	1	6	1	6.0
ZW03	3	4	7	1	7.0

A.7 Estimates of emissions from pneumatic controllers in the United States

National emissions are estimated by multiplying an average emission measurement by the number of times that emission occurs on the national scale. Emission measurements are often referred to as an “emission factor” or EF, and the data used to scale up the emissions is called the activity factor (AF). Emissions are calculated as:

$$EF_i * AF_i = ER_i \quad (A7.1)$$

where:

EF_i = Emission Factor for region i

AF_i = Activity Factor for region i

ER_i = resulting Emission Rate total for region i

For this work, the activity factors are national estimates of the number of pneumatic controllers. Emission factors are based on average emissions per controller measured in this work. Four combinations of activity factors and emission factors are used to calculate national emission estimates (Table A-12); these scenarios illustrate the uncertainty in national emission estimates that arise due to uncertainties in activity factors.

Table A-12. National methane emission estimates, assuming various activity factor scenarios

Activity Factor	Emission Factor	National methane emission estimate
Number of controllers in EPA 2012 Greenhouse Gas National Emission Inventory, 477,606 controllers	Average emission rate for all controllers measured in this work, 4.9 scf methane/h	394 Gg/y 20.5 bcf/y
Number of controllers in EPA 2012 Greenhouse Gas National Emission Inventory, 477,606 controllers	Average emission rate for controllers measured in this work, not including ESD controllers in the average, 5.5 scf methane/h	442 Gg/y 23.0 bcf/y
Number of wells in EPA 2012 Greenhouse Gas National Emission Inventory, 470,913 wells, with 75% of the wells assumed to have 2.7 pneumatic controllers per well, as observed in this work	Average emission rate for all controllers measured in this work, 4.9 scf methane/h	786 Gg/y 40.9 bcf/y
Number of wells in EPA 2012 Greenhouse Gas National Emission Inventory, 470,913 wells, with all of the wells assumed to have 2.7 pneumatic controllers per well, as observed in this work	Average emission rate for all controllers measured in this work, 4.9 scf methane/h	1050 Gg/y 54.6 bcf/y

The range of estimates (394-1,050 Gg/y) in Table S7-2 indicates the extent of variability in national emission estimates due to uncertainties in activity data. In the absence of additional activity data, a central estimate will be assumed to be a mid-point between the second and third scenarios, approximately 600 Gg/yr. These scenarios were selected since at least some of the participants in this study have significant numbers of wells with no pneumatic controllers, and because at least some of the participants do not inventory devices, such as Emergency Shut Down controllers, that are not expected to actuate.

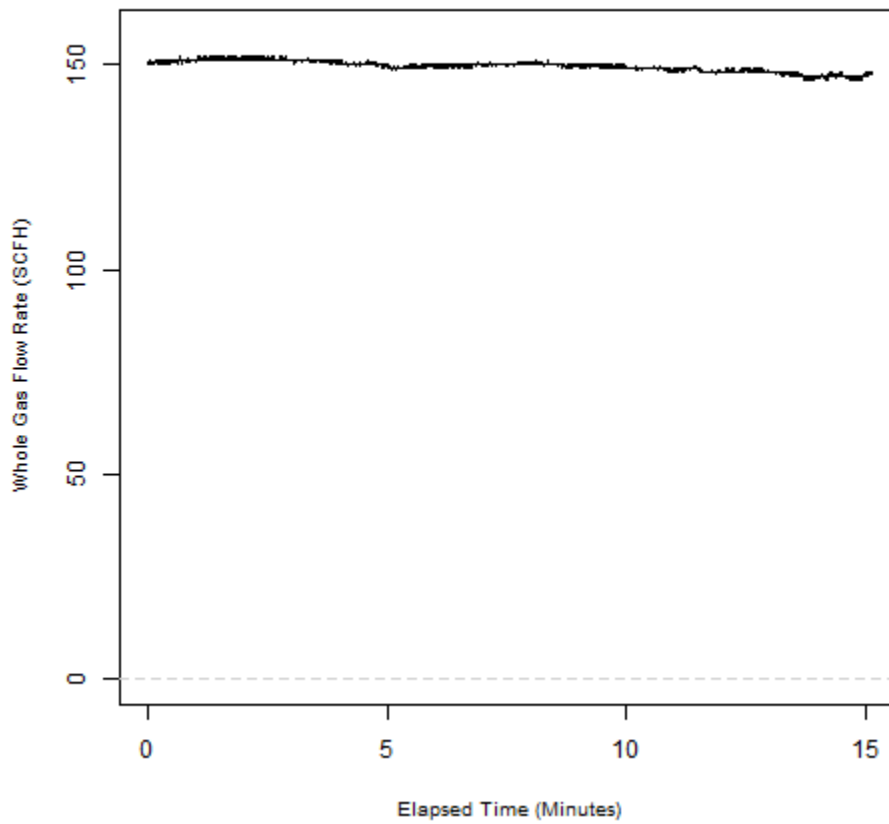
A.8 *Summary of data on 40 highest emitting devices*

Controller characteristics and time series for measured emissions are reported for the 40 devices with the highest measured emission rates. This information was reviewed by technical experts on pneumatic controller operation to assess whether controllers were operating properly. While these 40 controllers were not removed from operation and subjected to a rigorous performance analysis, the data that were collected may reveal some features of operation. For example, if a device had very high rates of nearly continuous emissions, larger than would be expected for the device design and supply gas pressure, it could be hypothesized that the controller/control valve system had a leak. If a device with intermittent actuation emissions, which appeared to be an intermittent vent controller, did not return to zero flow between actuations, it could be hypothesized that the barrier between the supply gas and the controller was not closing completely. It is important to recognize that these are hypotheses, based on expert opinion. The controllers are categorized, based on these hypotheses, as having “*equipment issues*” or “*operating as expected*”.

CZ09-PC01 [1] Region: GC

Device characteristics:

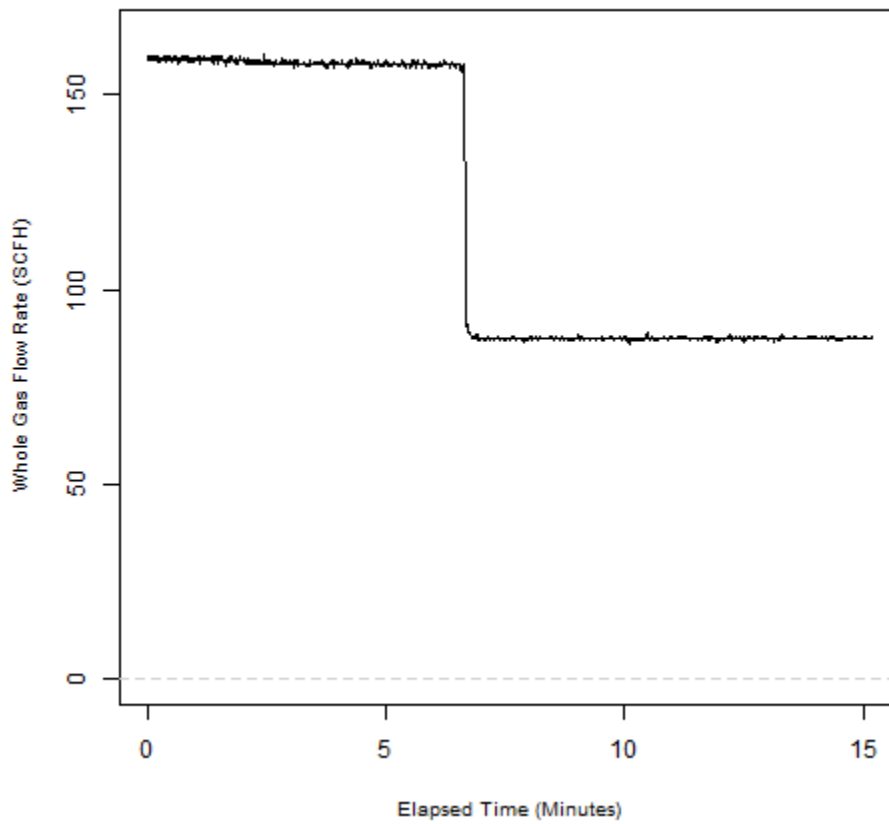
Avg. Emission rate (whole gas):	149.7 scf/h
Avg. Emission rate (methane):	143.8 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	19 psig
Assessment of controller operation:	equipment issues



CZ02-PC01 [2] Region: GC

Device characteristics:

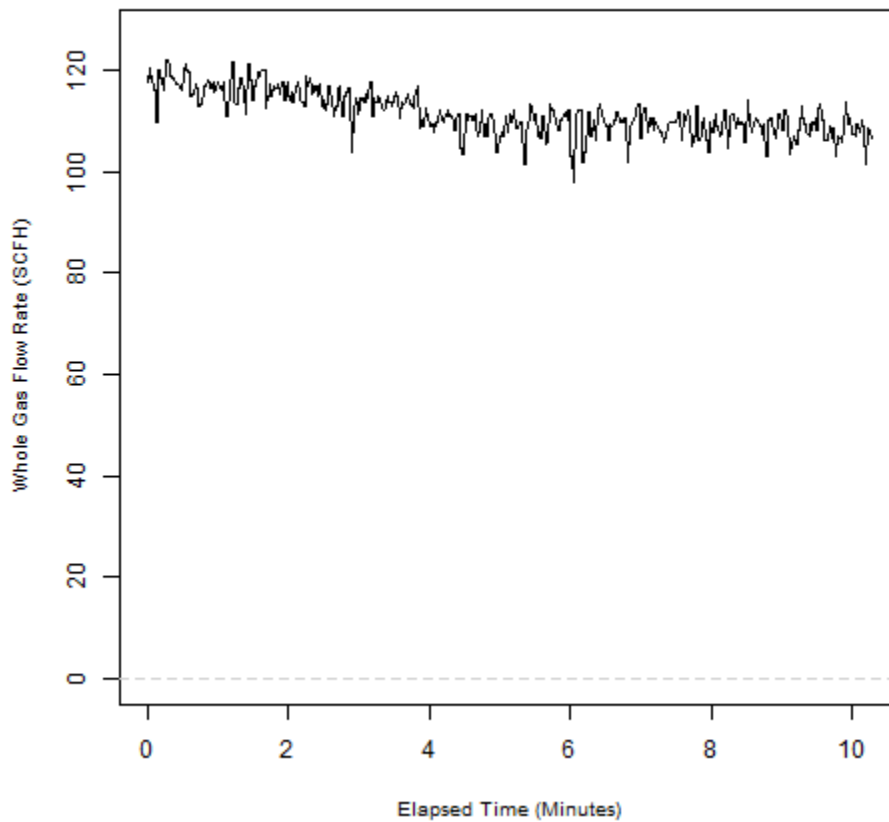
Avg. Emission rate (whole gas):	118.5 scf/h
Avg. Emission rate (methane):	108.0 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	J01
Supply pressure:	17 psig
Assessment of controller operation:	equipment issues



AA02-PC08 [3] Region: MC

Device characteristics:

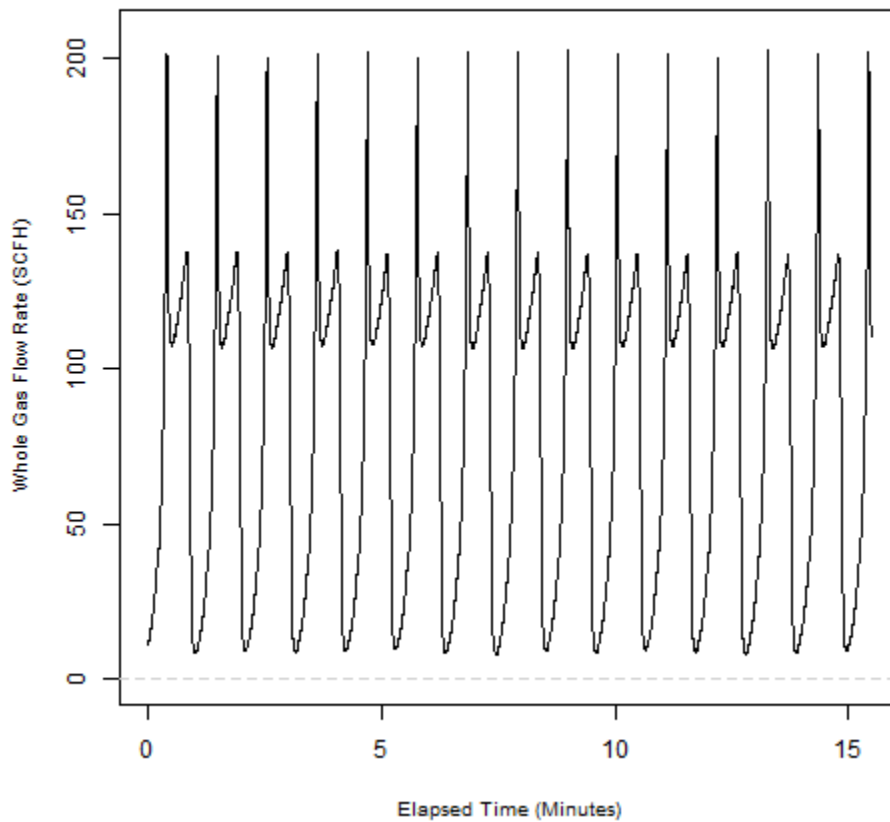
Avg. Emission rate (whole gas):	111.4 scf/h
Avg. Emission rate (methane):	89.7 scf/h
Controller application:	Pressure
Basic Application:	Compressor
Detailed Application:	Compressor - Suction Feed Control
Manufacturer/Model (blinded):	I01
Supply pressure:	60 psig
Assessment of controller operation:	equipment issues



LB05-PC03 [4] Region: GC

Device characteristics:

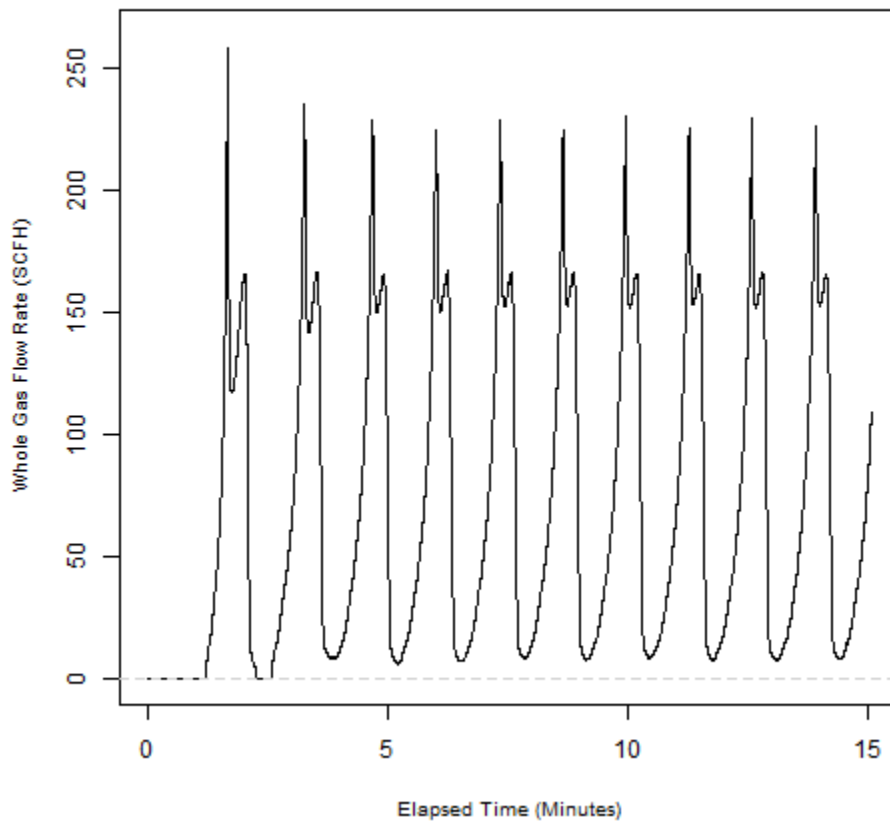
Avg. Emission rate (whole gas):	78.6 scf/h
Avg. Emission rate (methane):	68.9 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	M03
Supply pressure:	25 psig
Assessment of controller operation:	equipment issues



LB05-PC01 [5] Region: GC

Device characteristics:

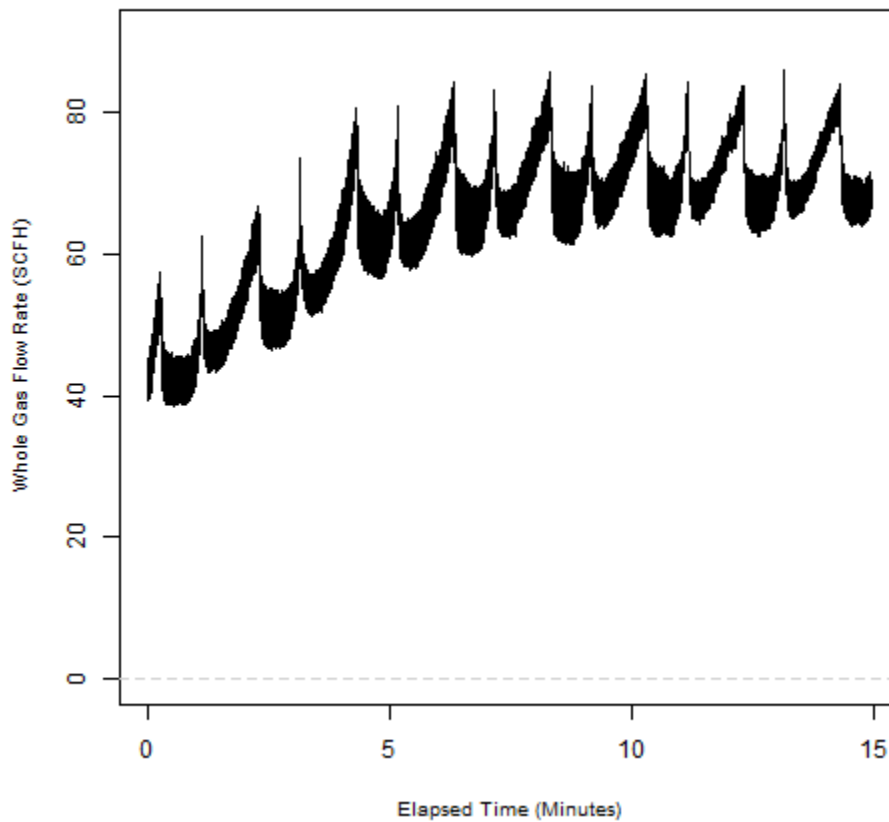
Avg. Emission rate (whole gas):	68.6 scf/h
Avg. Emission rate (methane):	60.2 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	M03
Supply pressure:	30 psig
Assessment of controller operation:	equipment issues



LB07-PC02 [6] Region: GC

Device characteristics:

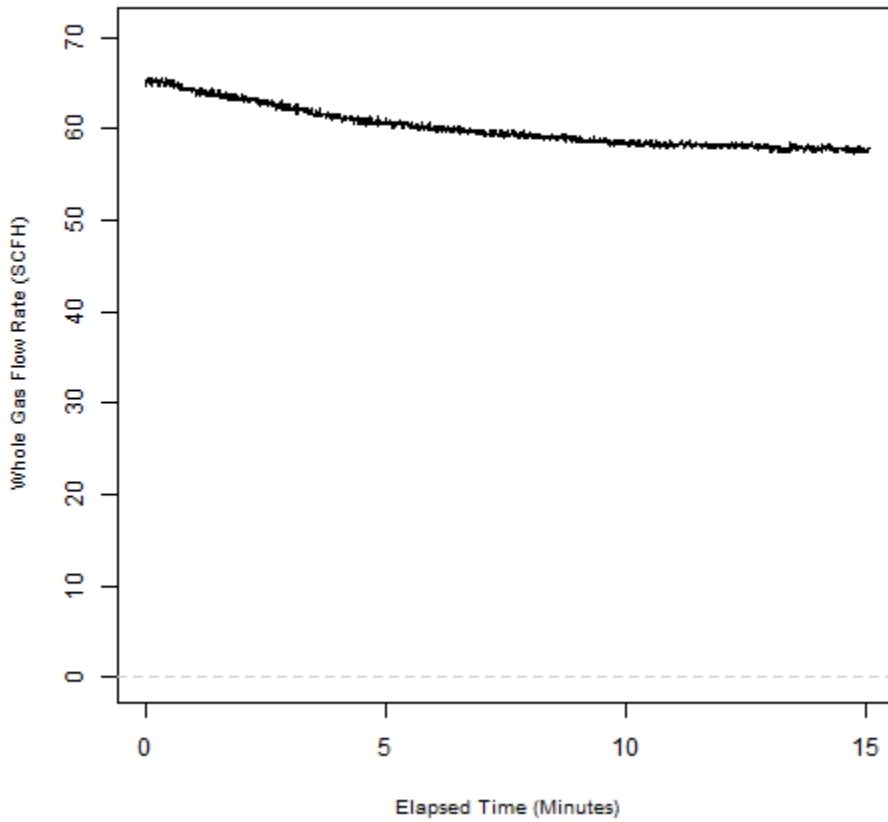
Avg. Emission rate (whole gas):	64.8 scf/h
Avg. Emission rate (methane):	51.2 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	M03
Supply pressure:	24 psig
Assessment of controller operation:	equipment issues



CZ08-PC01 [7] Region: GC

Device characteristics:

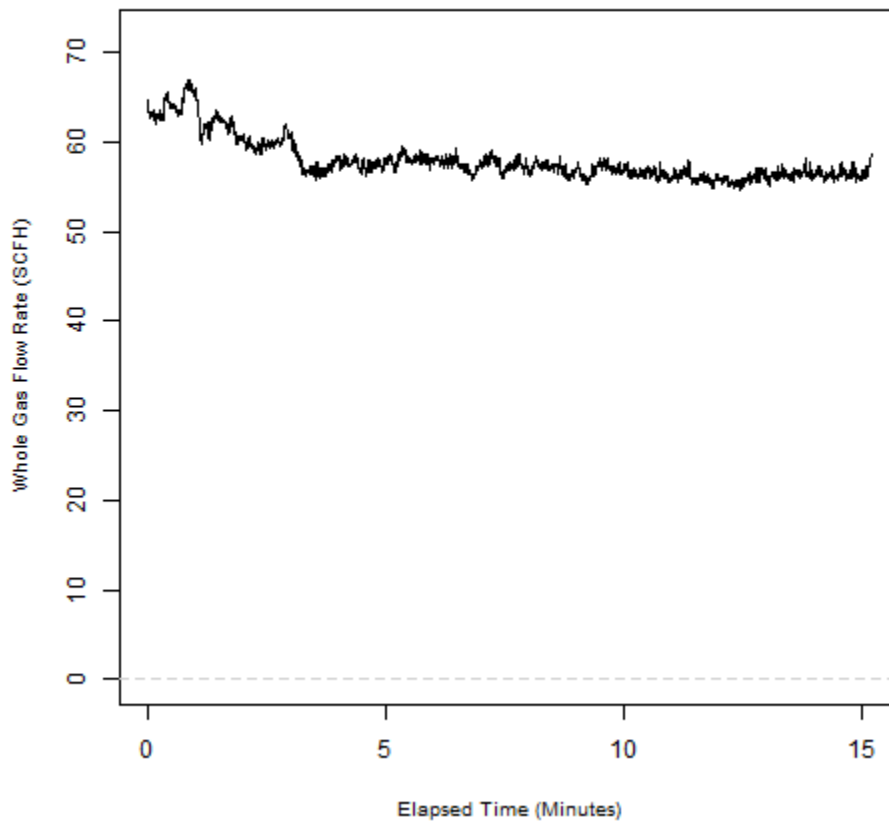
Avg. Emission rate (whole gas):	60.1 scf/h
Avg. Emission rate (methane):	57.8 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	11 psig
Assessment of controller operation:	equipment issues



GZ02-PC12 [8] Region: GC

Device characteristics:

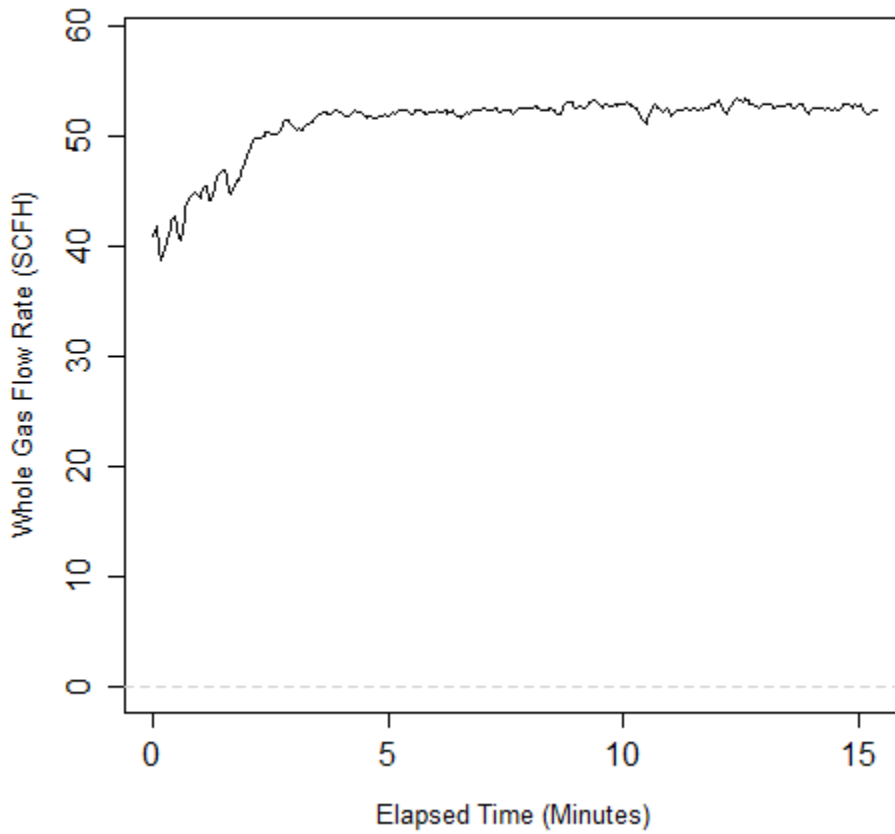
Avg. Emission rate (whole gas):	57.9 scf/h
Avg. Emission rate (methane):	56.5 scf/h
Controller application:	Level
Basic Application:	Compressor
Detailed Application:	Compressor - Liquid Level Control (Stage 2)
Manufacturer/Model (blinded):	J01
Supply pressure:	28 psig
Assessment of controller operation:	equipment issues



CZ06-PC01 [9] Region: GC

Device characteristics:

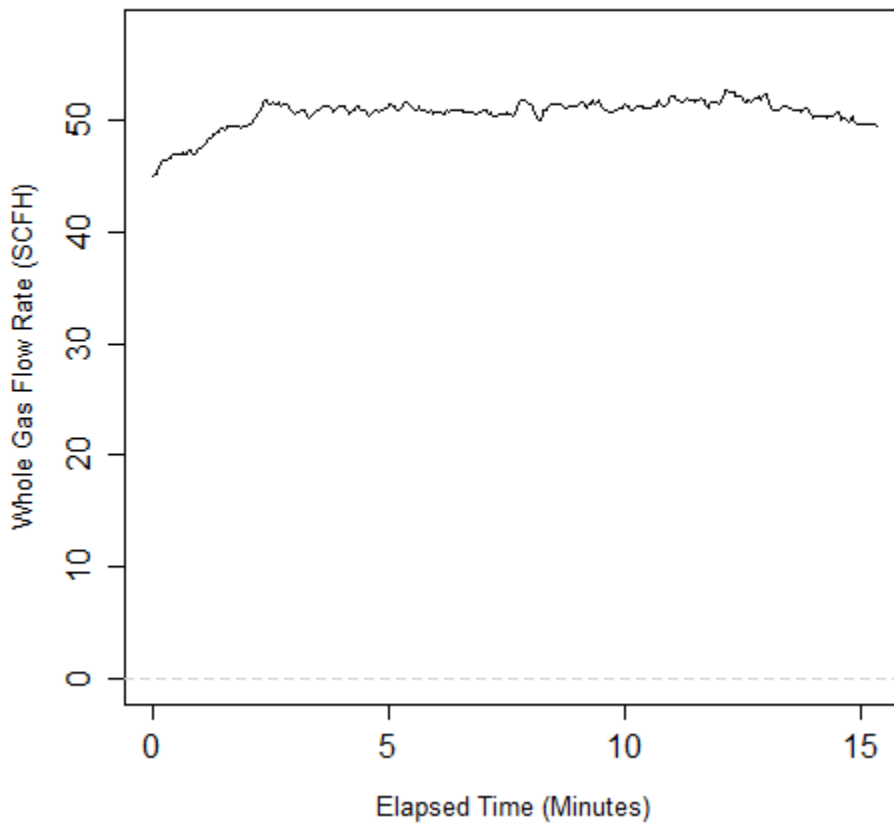
Avg. Emission rate (whole gas):	51.2 scf/h
Avg. Emission rate (methane):	48.2 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	G01
Supply pressure:	6 psig
Assessment of controller operation:	operating as expected



CZ06-PC02 [10] Region: GC

Device characteristics:

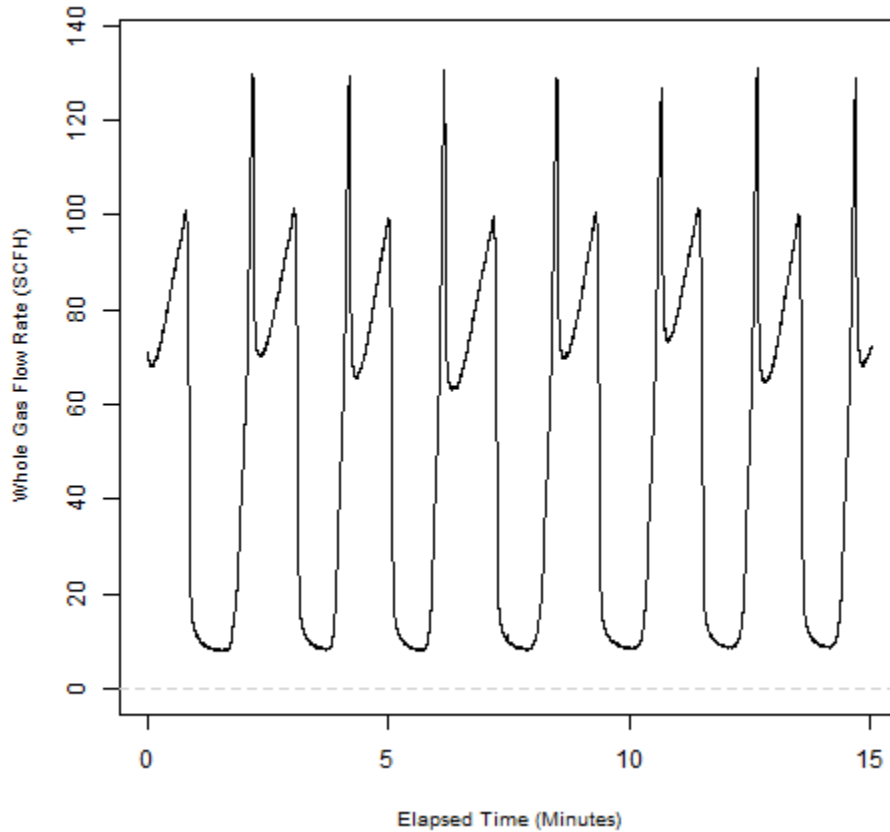
Avg. Emission rate (whole gas):	50.7 scf/h
Avg. Emission rate (methane):	47.7 scf/h
Controller application:	Pressure
Basic Application:	Plunger Lift
Detailed Application:	Plunger Lift
Manufacturer/Model (blinded):	E01
Supply pressure:	20 psig
Assessment of controller operation:	equipment issues



LB03-PC01 [11] Region: GC

Device characteristics:

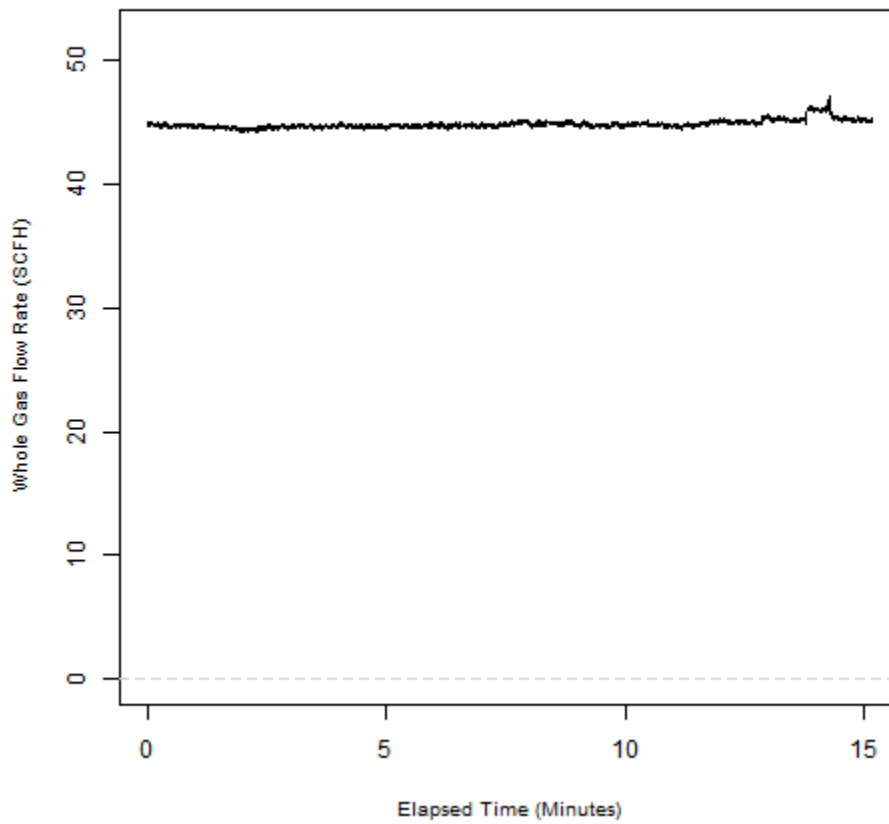
Avg. Emission rate (whole gas):	50.0 scf/h
Avg. Emission rate (methane):	39.7 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	30 psig
Assessment of controller operation:	equipment issues



CZ09-PC02 [12] Region: GC

Device characteristics:

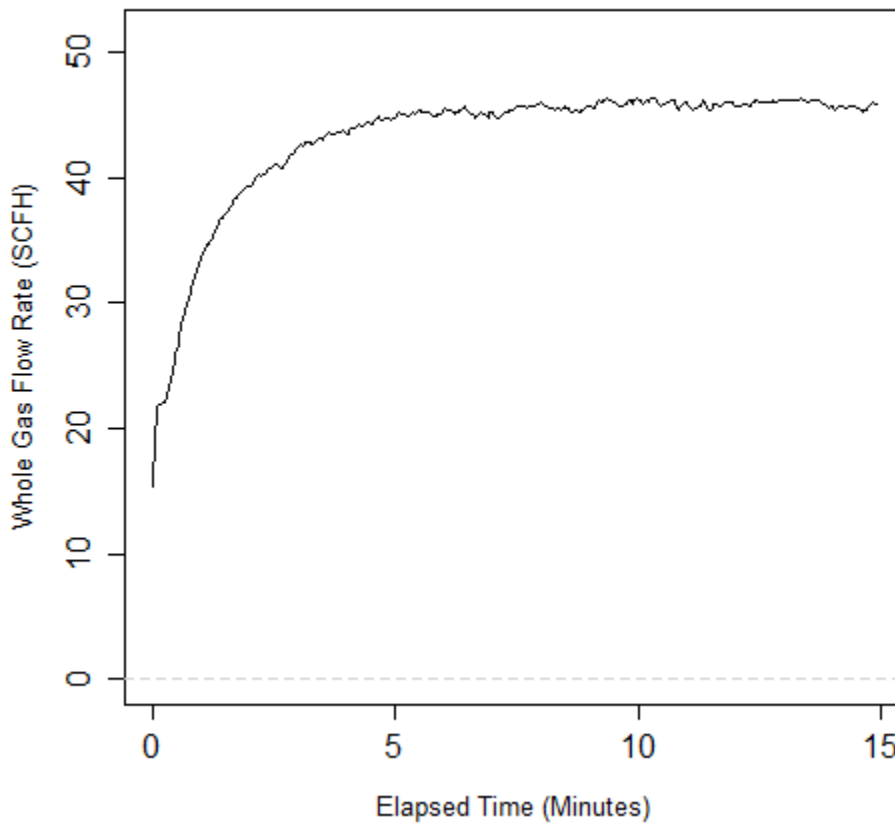
Avg. Emission rate (whole gas):	44.9 scf/h
Avg. Emission rate (methane):	43.1 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	9 psig
Assessment of controller operation:	equipment issues



GZ02-PC14 [13] Region: GC

Device characteristics:

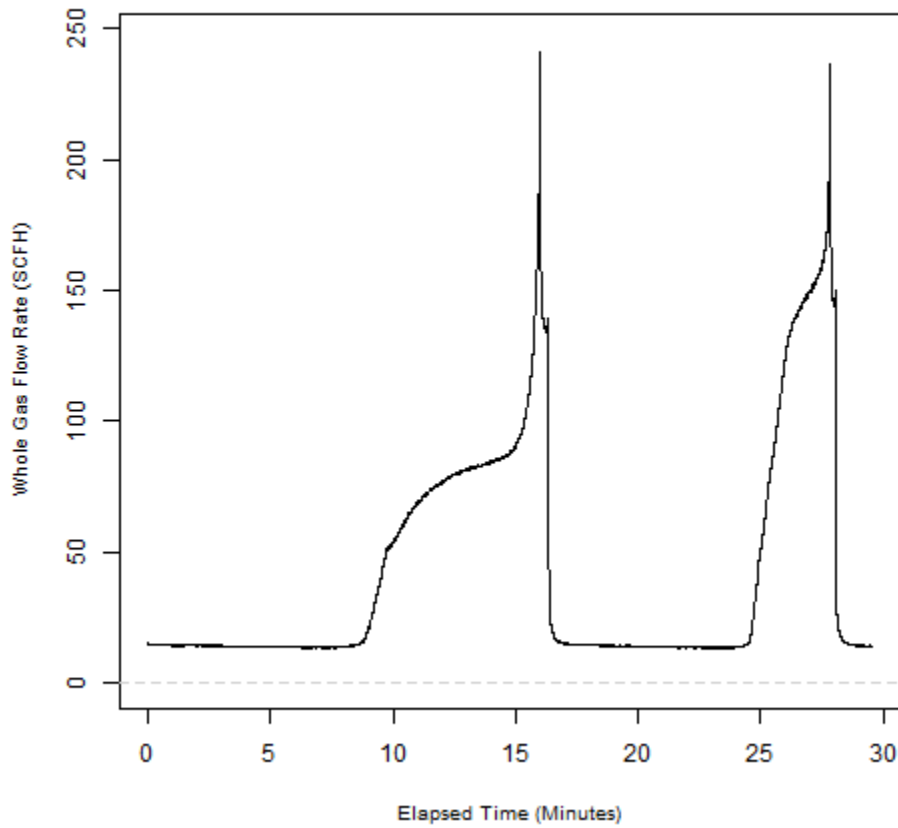
Avg. Emission rate (whole gas):	43.3 scf/h
Avg. Emission rate (methane):	42.2 scf/h
Controller application:	Pressure
Basic Application:	Compressor
Detailed Application:	Compressor - Suction Feed Control
Manufacturer/Model (blinded):	I01
Supply pressure:	42 psig
Assessment of controller operation:	equipment issues



CZ10-PC01 [14] Region: GC

Device characteristics:

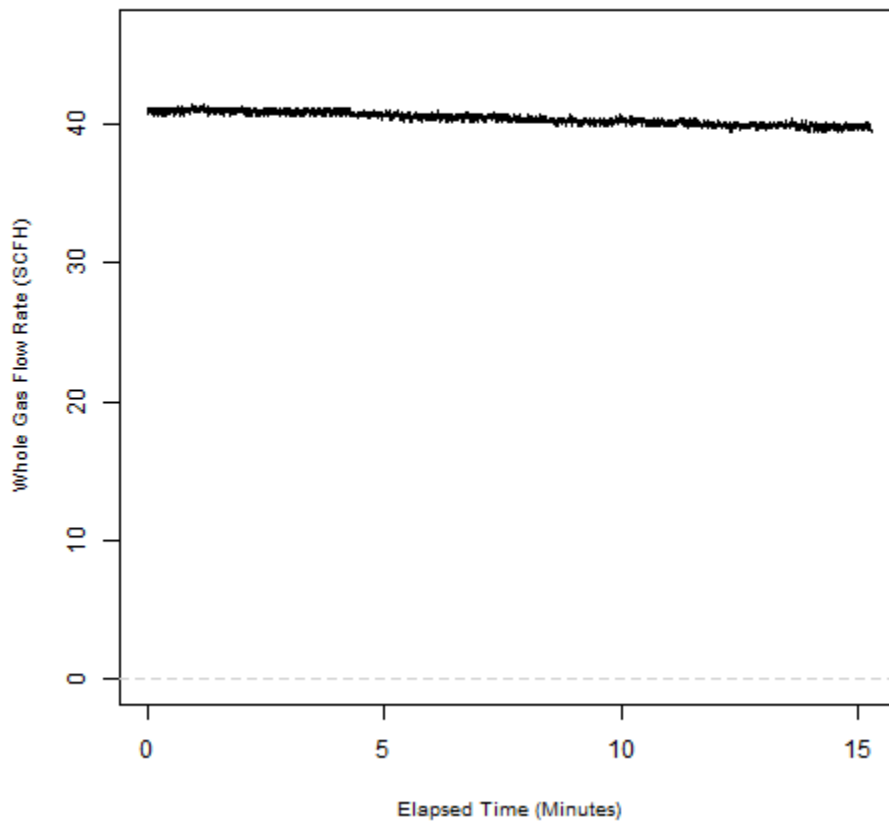
Avg. Emission rate (whole gas):	43.2 scf/h
Avg. Emission rate (methane):	41.5 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	13 psig
Assessment of controller operation:	equipment issues



CZ05-PC01 [15] Region: GC

Device characteristics:

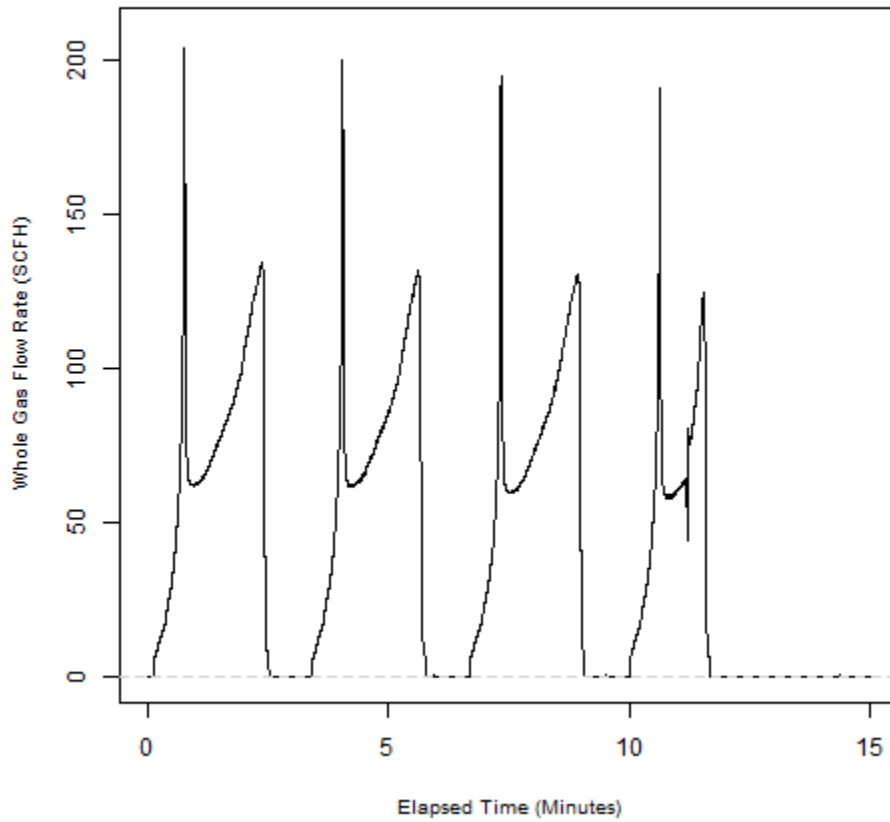
Avg. Emission rate (whole gas):	40.4 scf/h
Avg. Emission rate (methane):	37.9 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	A01
Supply pressure:	29 psig
Assessment of controller operation:	equipment issues



LB01-PC01 [16] Region: GC

Device characteristics:

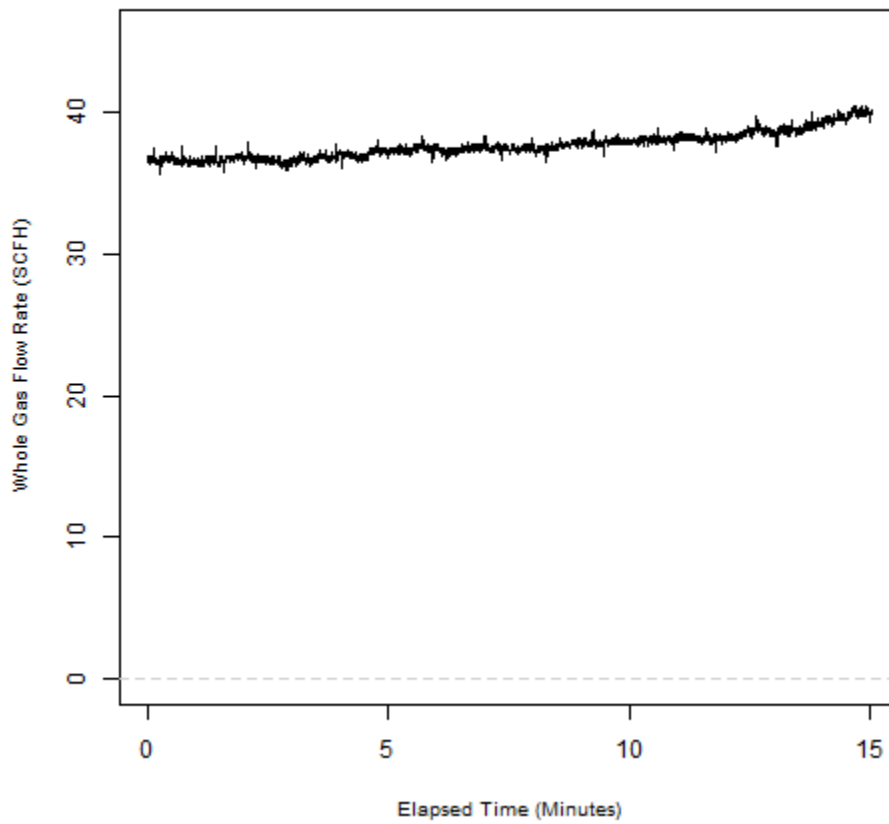
Avg. Emission rate (whole gas):	40.2 scf/h
Avg. Emission rate (methane):	32.3 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	M03
Supply pressure:	30 psig
Assessment of controller operation:	operating as expected



GZ03-PC22 [17] Region: GC

Device characteristics:

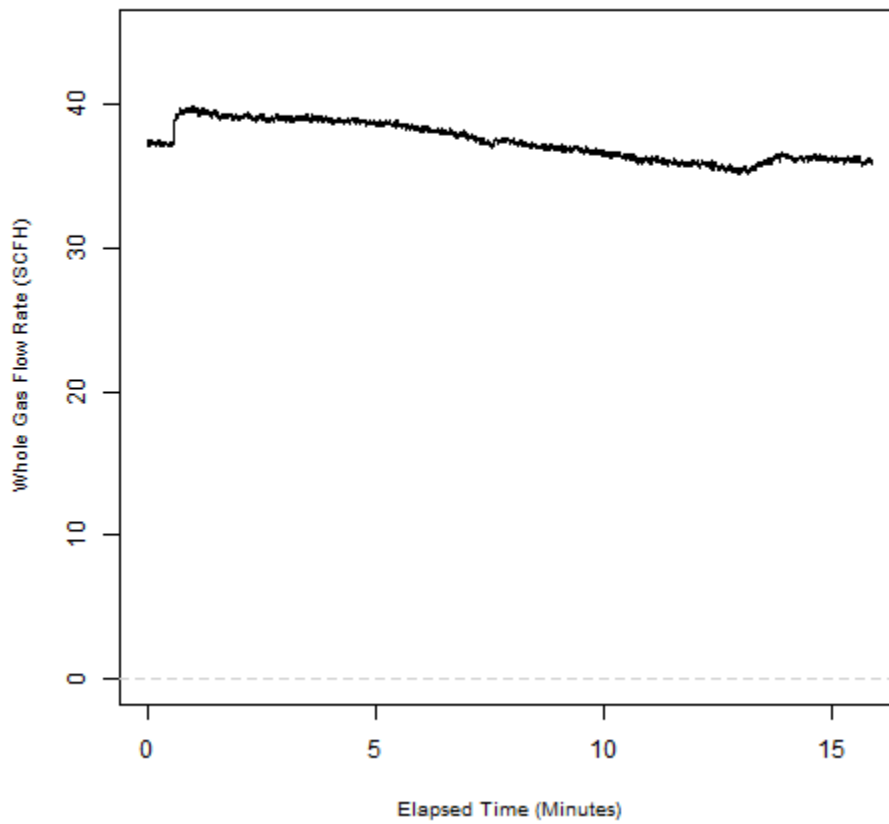
Avg. Emission rate (whole gas):	37.7 scf/h
Avg. Emission rate (methane):	36.6 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	7 psig
Assessment of controller operation:	equipment issues



XQ04-PC02 [18] Region: MC

Device characteristics:

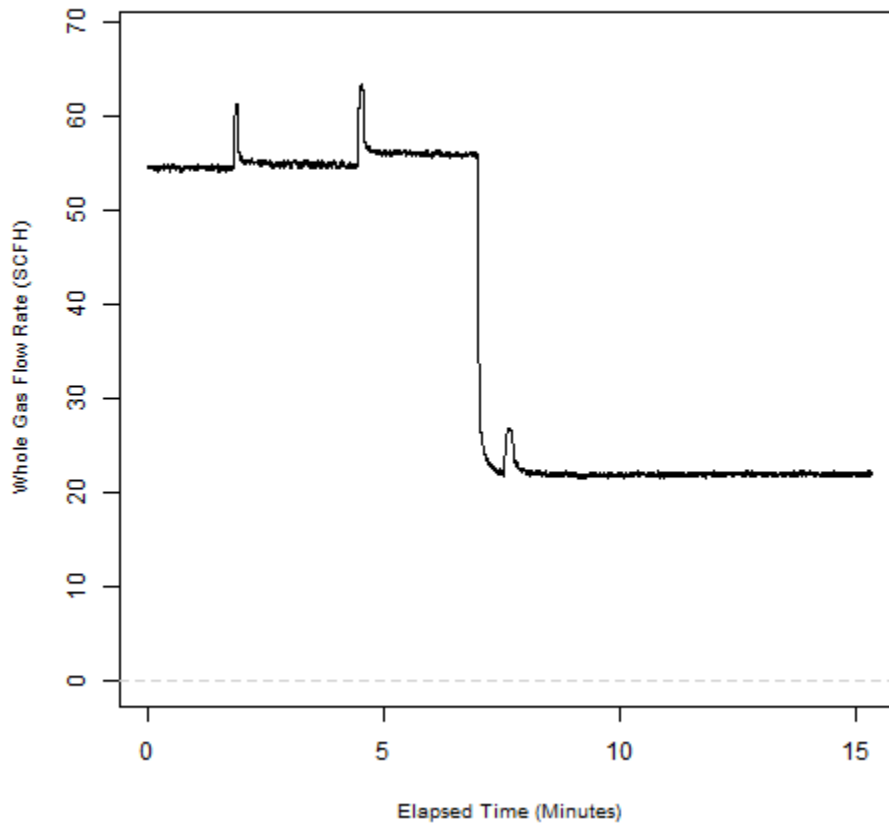
Avg. Emission rate (whole gas):	37.4 scf/h
Avg. Emission rate (methane):	33.5 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	Q02
Supply pressure:	28 psig
Assessment of controller operation:	equipment issues



XQ01-PC04 [19] Region: MC

Device characteristics:

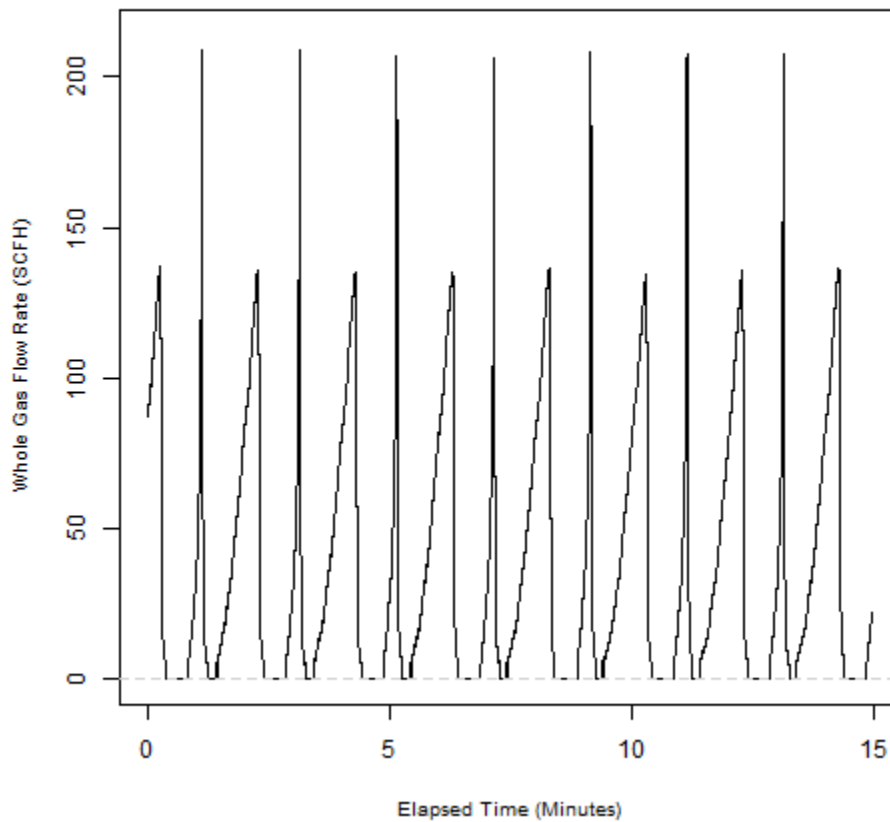
Avg. Emission rate (whole gas):	37.4 scf/h
Avg. Emission rate (methane):	33.0 scf/h
Controller application:	Pressure
Basic Application:	Plunger Lift
Detailed Application:	Plunger Lift
Manufacturer/Model (blinded):	F02
Supply pressure:	15 psig
Assessment of controller operation:	equipment issues



LB07-PC01 [20] Region: GC

Device characteristics:

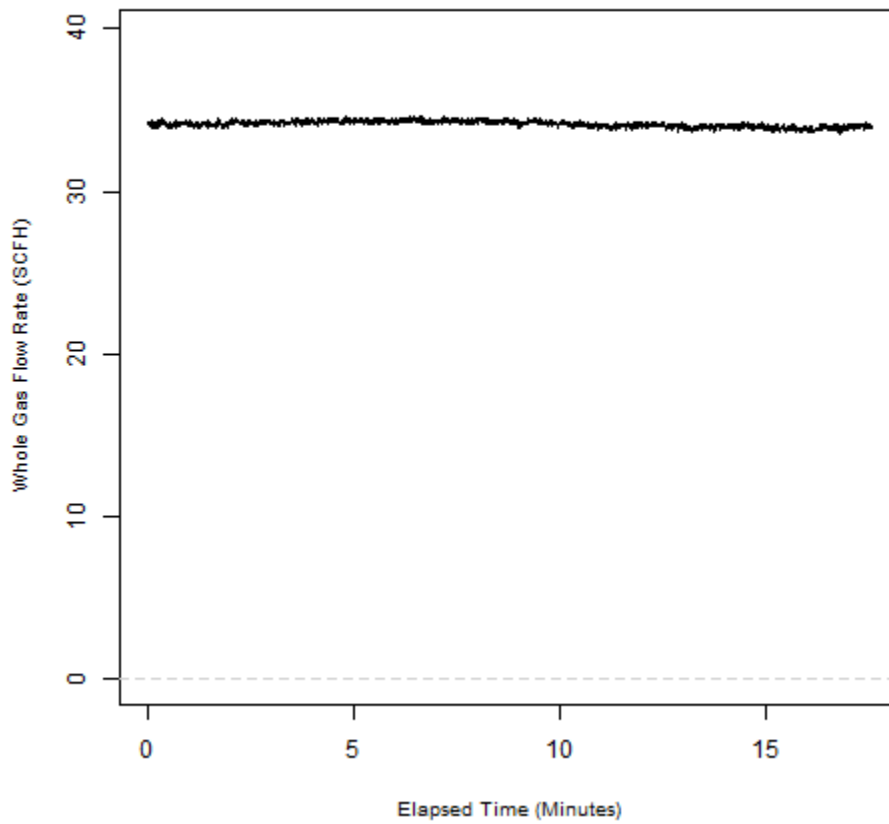
Avg. Emission rate (whole gas):	36.3 scf/h
Avg. Emission rate (methane):	28.7 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	M03
Supply pressure:	24 psig
Assessment of controller operation:	operating as expected



GZ03-PC13 [21] Region: GC

Device characteristics:

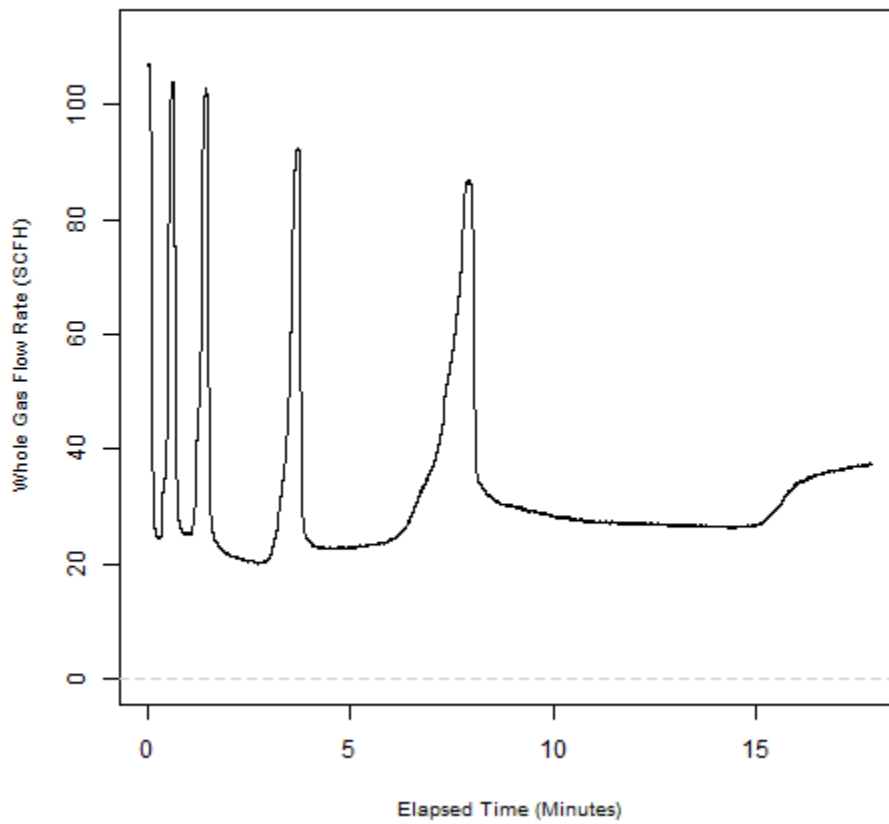
Avg. Emission rate (whole gas):	34.1 scf/h
Avg. Emission rate (methane):	33.2 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	26 psig
Assessment of controller operation:	equipment issues



GZ04-PC03 [22] Region: GC

Device characteristics:

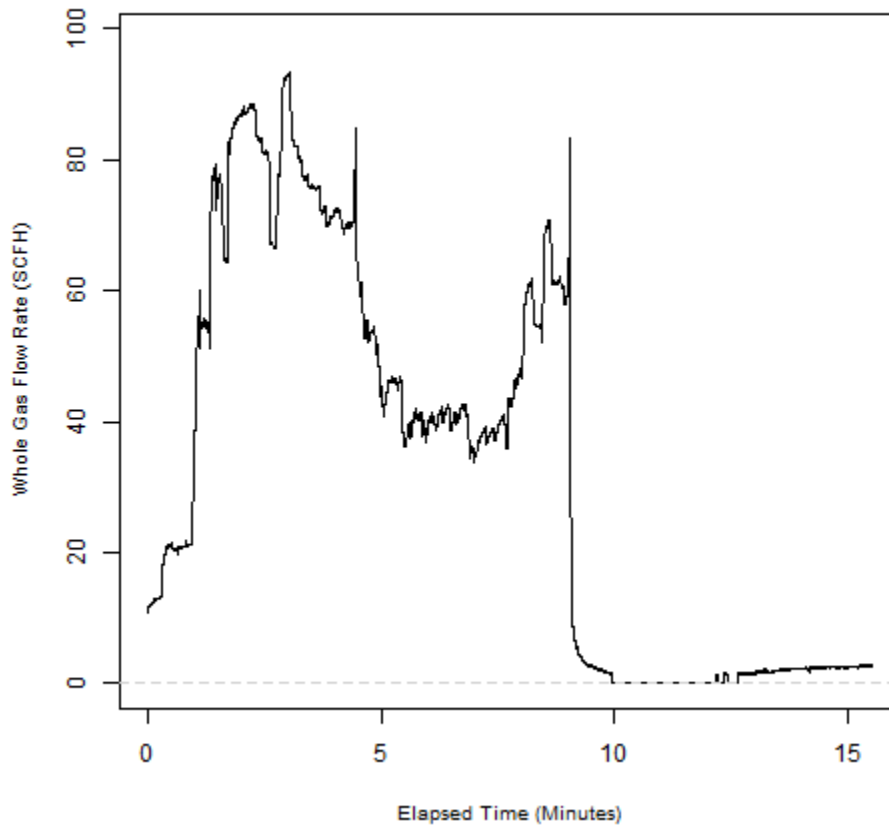
Avg. Emission rate (whole gas):	32.7 scf/h
Avg. Emission rate (methane):	31.8 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	30 psig
Assessment of controller operation:	equipment issues



GZ03-PC02 [23] Region: GC

Device characteristics:

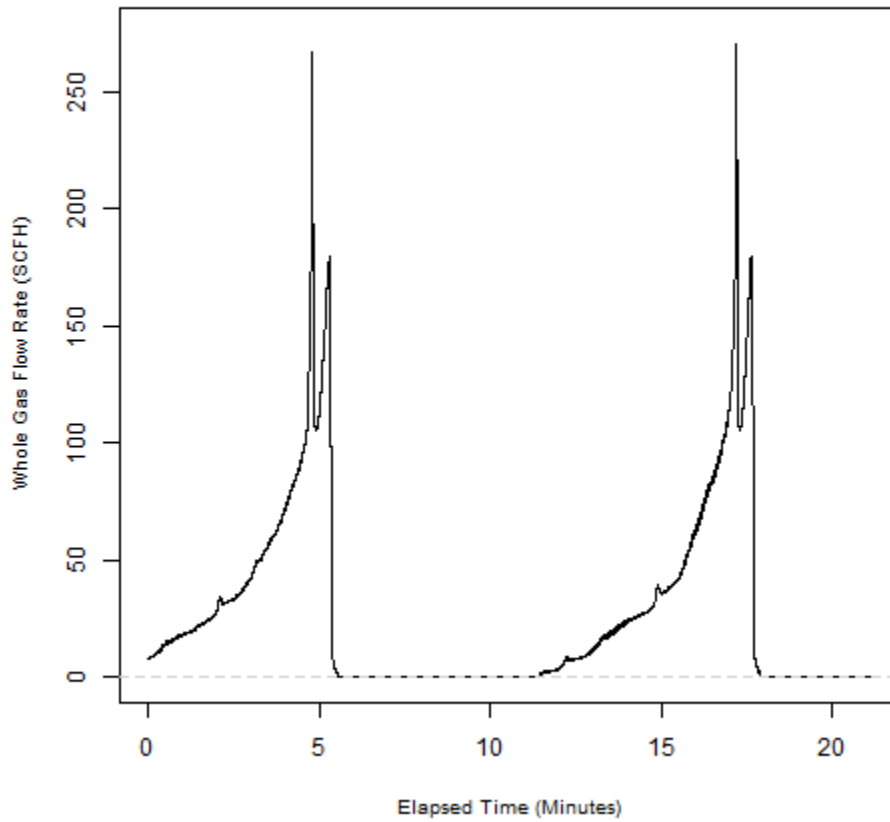
Avg. Emission rate (whole gas):	32.6 scf/h
Avg. Emission rate (methane):	31.7 scf/h
Controller application:	Level
Basic Application:	Compressor
Detailed Application:	Compressor - Liquid Level Control (Stage 2)
Manufacturer/Model (blinded):	L01
Supply pressure:	30 psig
Assessment of controller operation:	operating as expected



LB07-PC04 [24] Region: GC

Device characteristics:

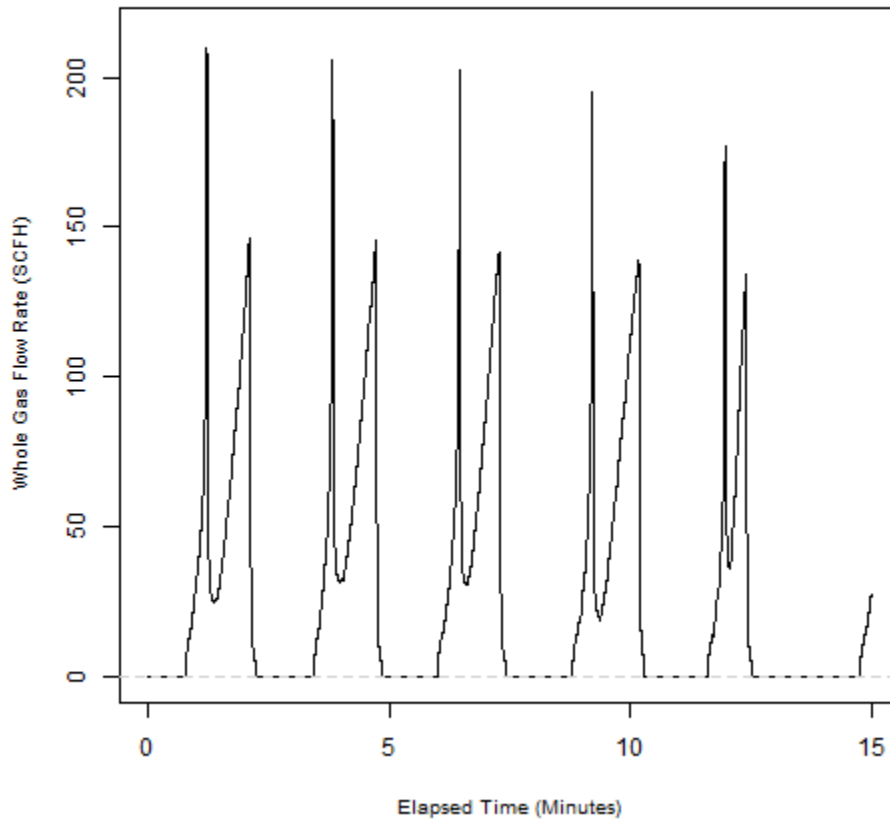
Avg. Emission rate (whole gas):	27.0 scf/h
Avg. Emission rate (methane):	21.4 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	M03
Supply pressure:	28 psig
Assessment of controller operation:	equipment issues



LB04-PC01 [25] Region: GC

Device characteristics:

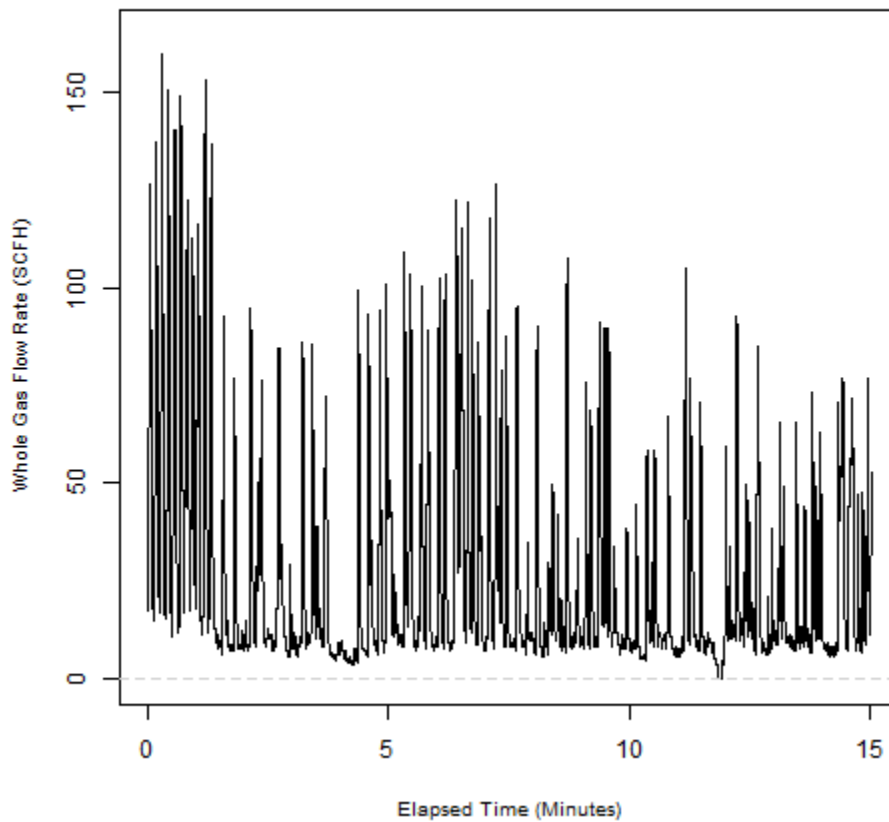
Avg. Emission rate (whole gas):	26.2 scf/h
Avg. Emission rate (methane):	20.7 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	M03
Supply pressure:	30 psig
Assessment of controller operation:	operating as expected



AP01-PC12 [26] Region: MC

Device characteristics:

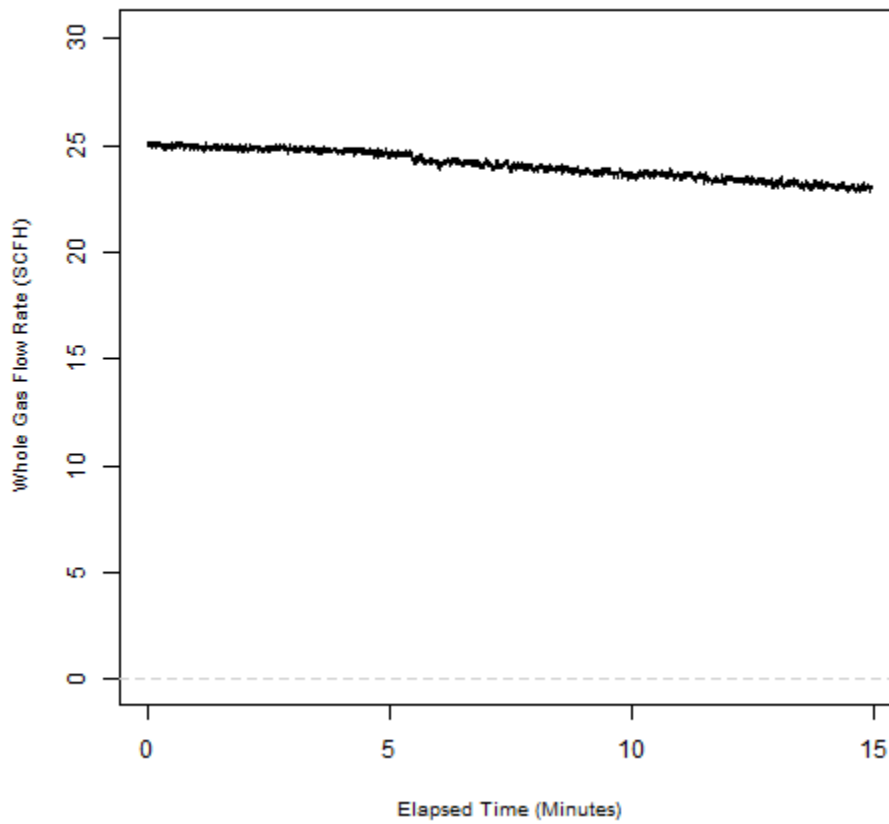
Avg. Emission rate (whole gas):	25.6 scf/h
Avg. Emission rate (methane):	17.0 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	I02
Supply pressure:	11 psig
Assessment of controller operation:	equipment issues



XQ05-PC02 [27] Region: MC

Device characteristics:

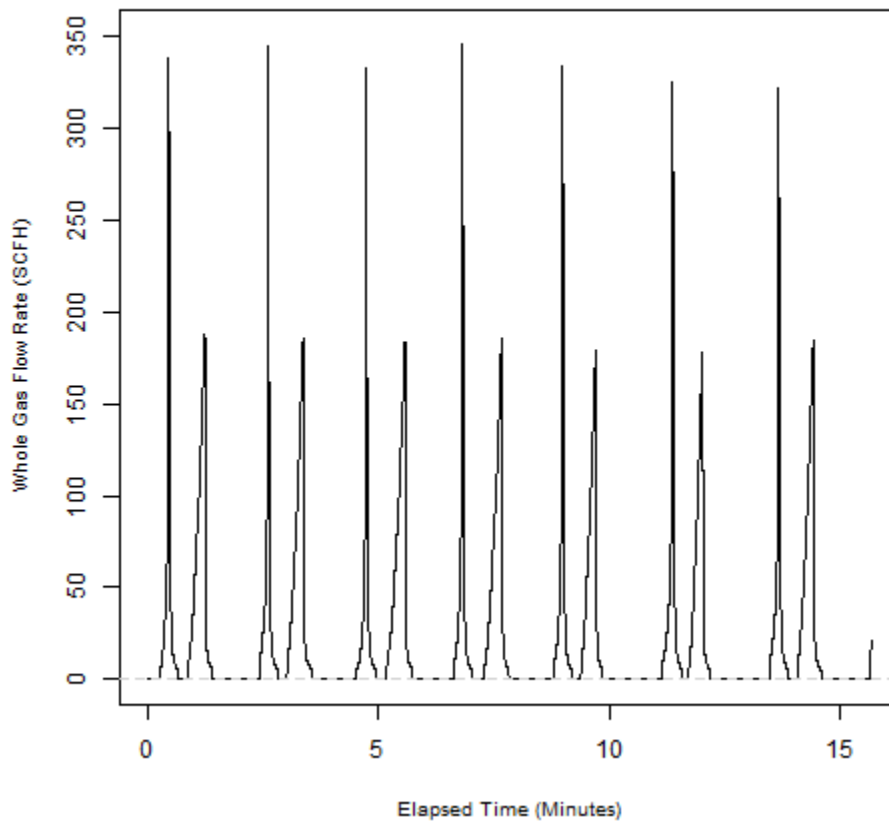
Avg. Emission rate (whole gas):	24.1 scf/h
Avg. Emission rate (methane):	21.7 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	Q02
Supply pressure:	27 psig
Assessment of controller operation:	operating as expected



LB06-PC05 [28] Region: GC

Device characteristics:

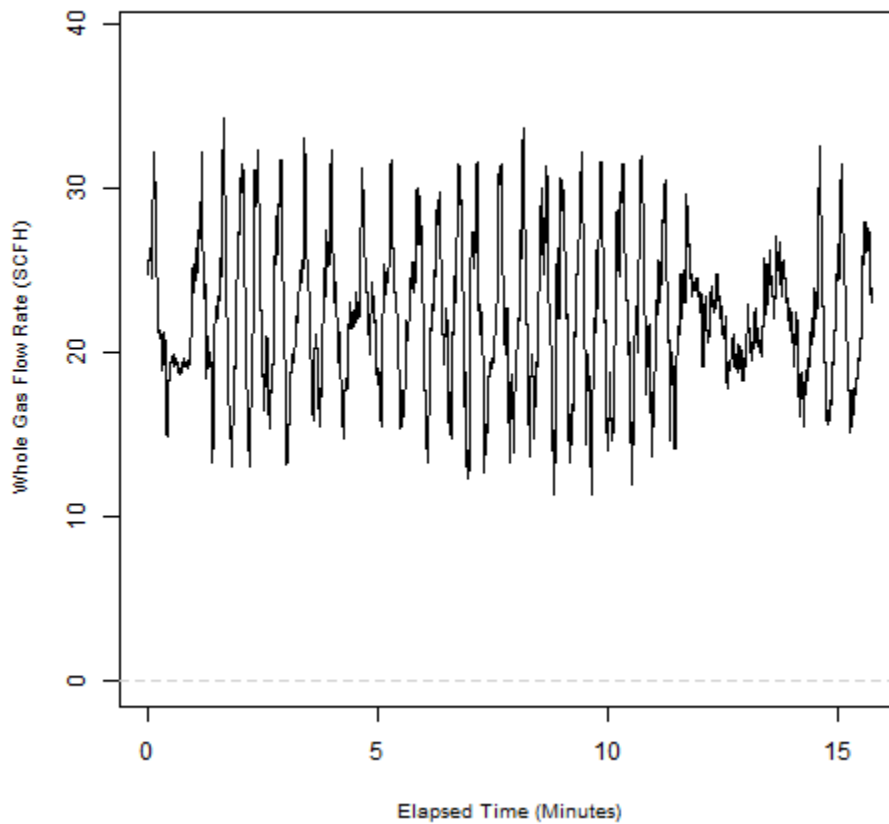
Avg. Emission rate (whole gas):	22.5 scf/h
Avg. Emission rate (methane):	18.3 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	M03
Supply pressure:	30 psig
Assessment of controller operation:	operating as expected



CZ11-PC01 [29] Region: GC

Device characteristics:

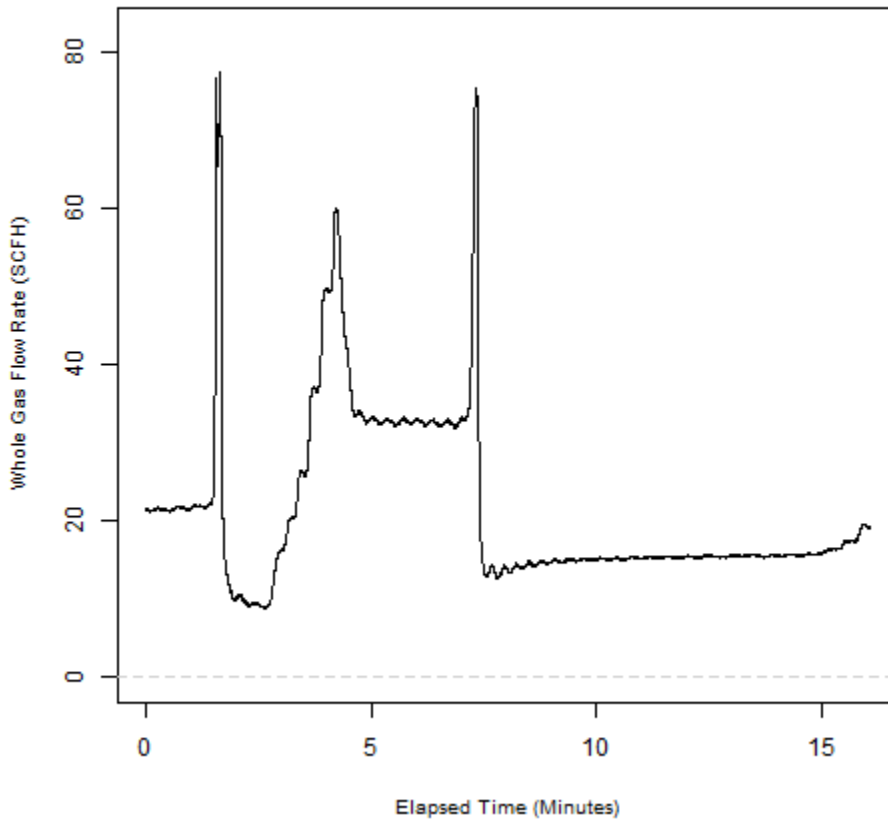
Avg. Emission rate (whole gas):	22.2 scf/h
Avg. Emission rate (methane):	21.3 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	23 psig
Assessment of controller operation:	equipment issues



CZ08-PC02 [30] Region: GC

Device characteristics:

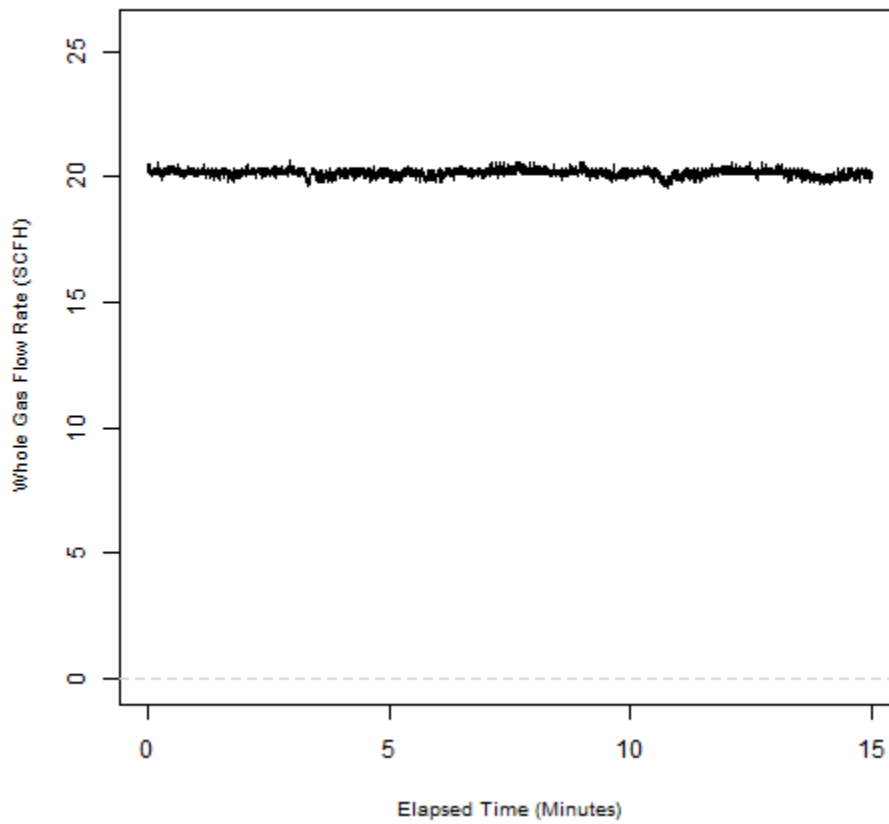
Avg. Emission rate (whole gas):	21.6 scf/h
Avg. Emission rate (methane):	21.1 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	19 psig
Assessment of controller operation:	equipment issues



XQ06-PC01 [31] Region: MC

Device characteristics:

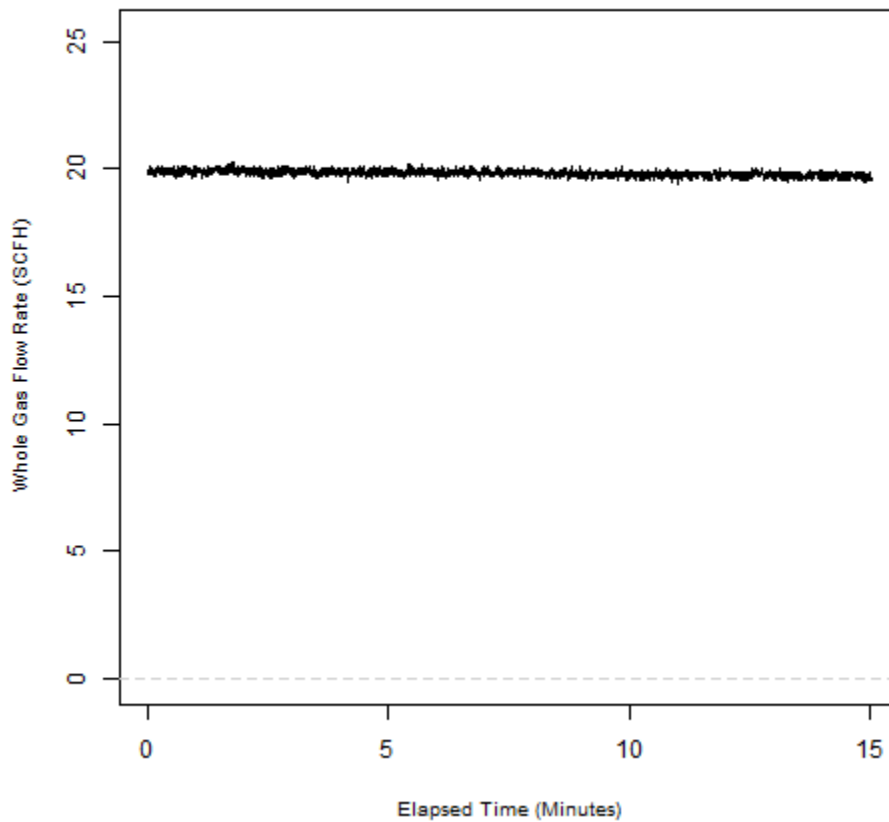
Avg. Emission rate (whole gas):	20.2 scf/h
Avg. Emission rate (methane):	18.0 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	G01
Supply pressure:	28 psig
Assessment of controller operation:	operating as expected



RB01-PC23 [32] Region: GC

Device characteristics:

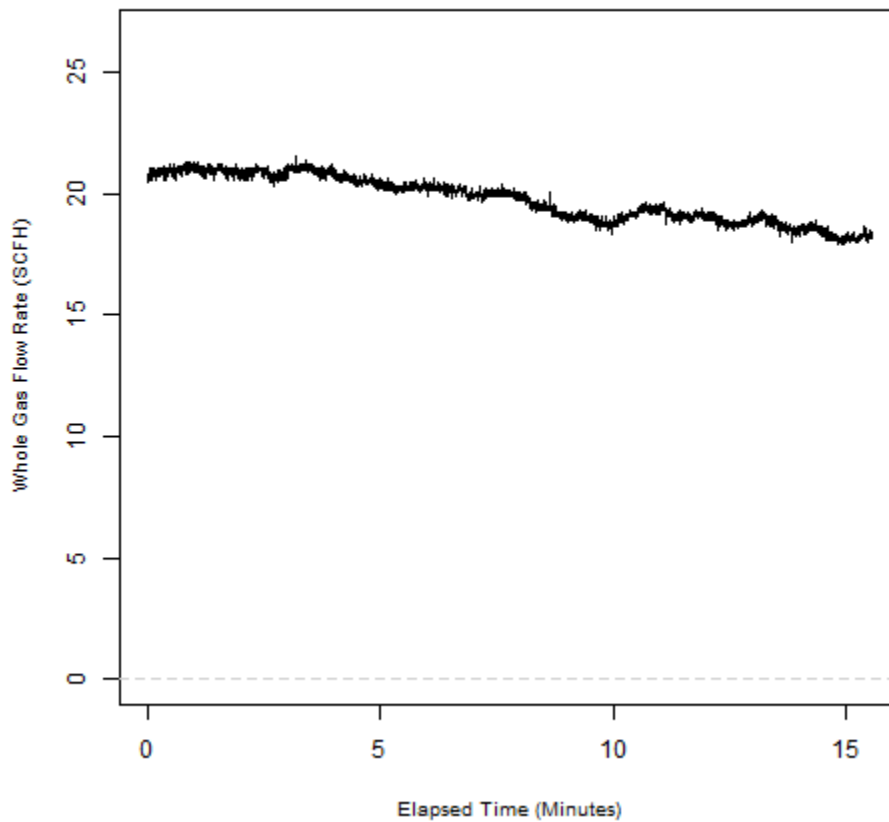
Avg. Emission rate (whole gas):	19.8 scf/h
Avg. Emission rate (methane):	16.3 scf/h
Controller application:	Pressure
Basic Application:	Dehydration System
Detailed Application:	Site Gas Feed Control
Manufacturer/Model (blinded):	M05
Supply pressure:	30 psig
Assessment of controller operation:	operating as expected



DL01-PC24 [33] Region: RM

Device characteristics:

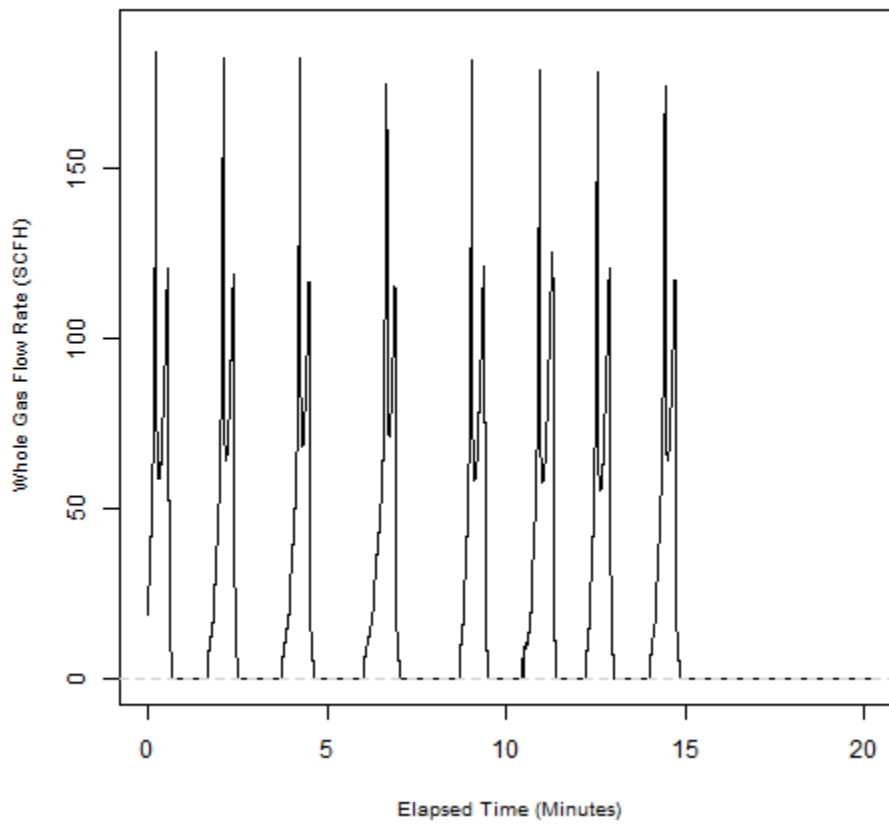
Avg. Emission rate (whole gas):	19.8 scf/h
Avg. Emission rate (methane):	17.7 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Liquid Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	38 psig
Assessment of controller operation:	equipment issues



LB04-PC03 [34] Region: GC

Device characteristics:

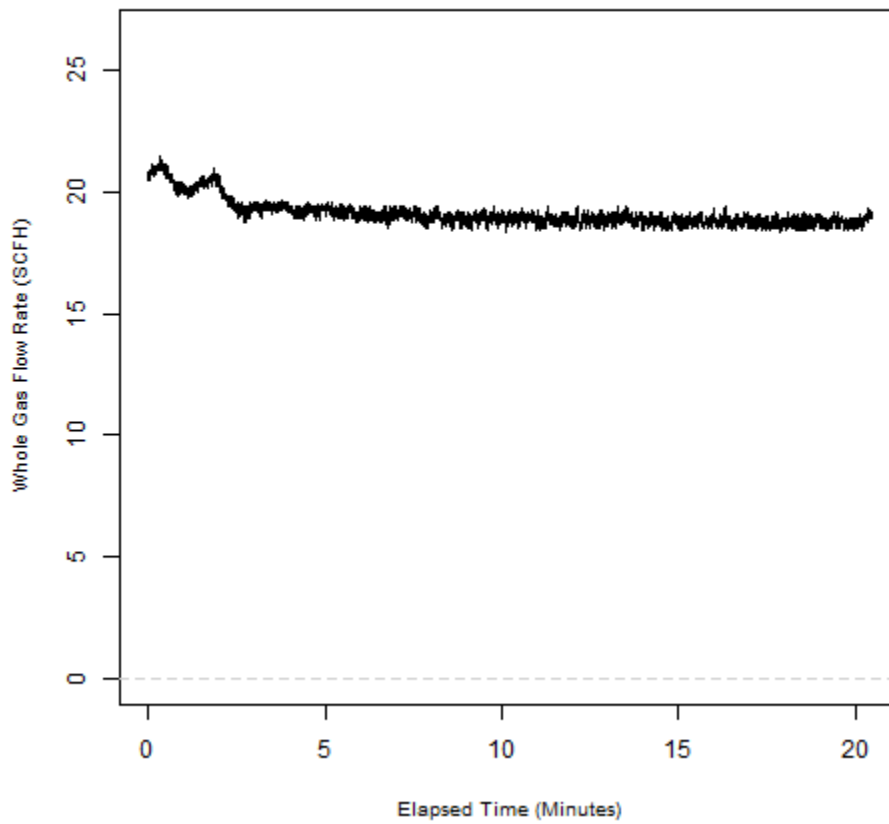
Avg. Emission rate (whole gas):	19.1 scf/h
Avg. Emission rate (methane):	15.0 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	M03
Supply pressure:	30 psig
Assessment of controller operation:	operating as expected



RQ03-PC01 [35] Region: MC

Device characteristics:

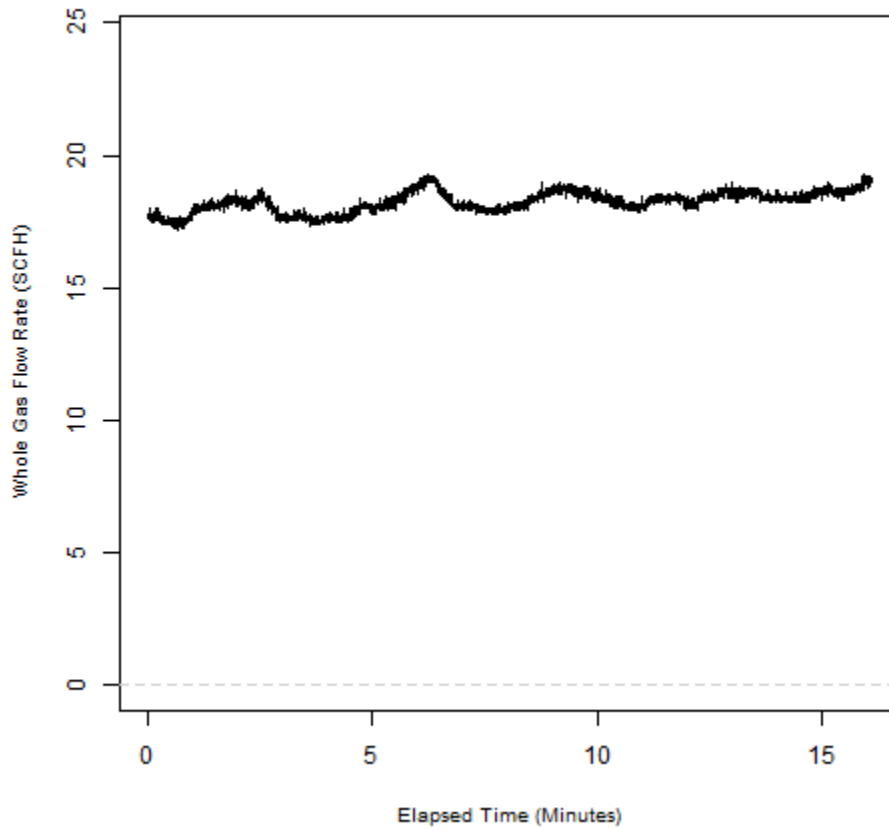
Avg. Emission rate (whole gas):	19.1 scf/h
Avg. Emission rate (methane):	16.3 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	Q02
Supply pressure:	22 psig
Assessment of controller operation:	operating as expected



RQ07-PC03 [36] Region: MC

Device characteristics:

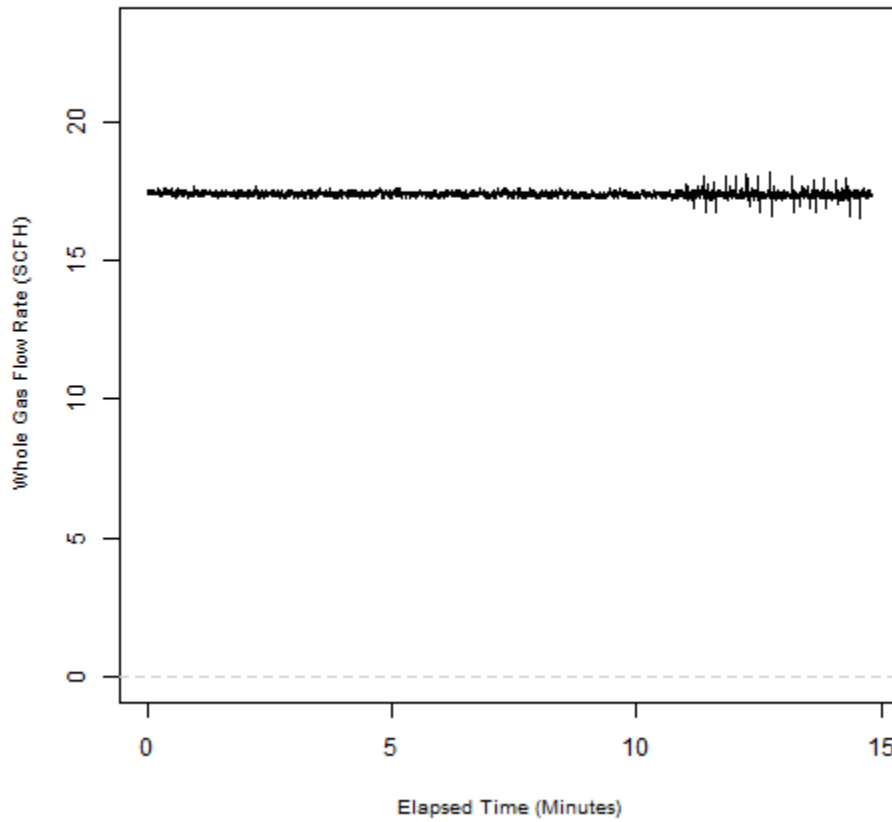
Avg. Emission rate (whole gas):	18.2 scf/h
Avg. Emission rate (methane):	14.2 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	G01
Supply pressure:	22 psig
Assessment of controller operation:	operating as expected



RQ05-PC03 [37] Region: MC

Device characteristics:

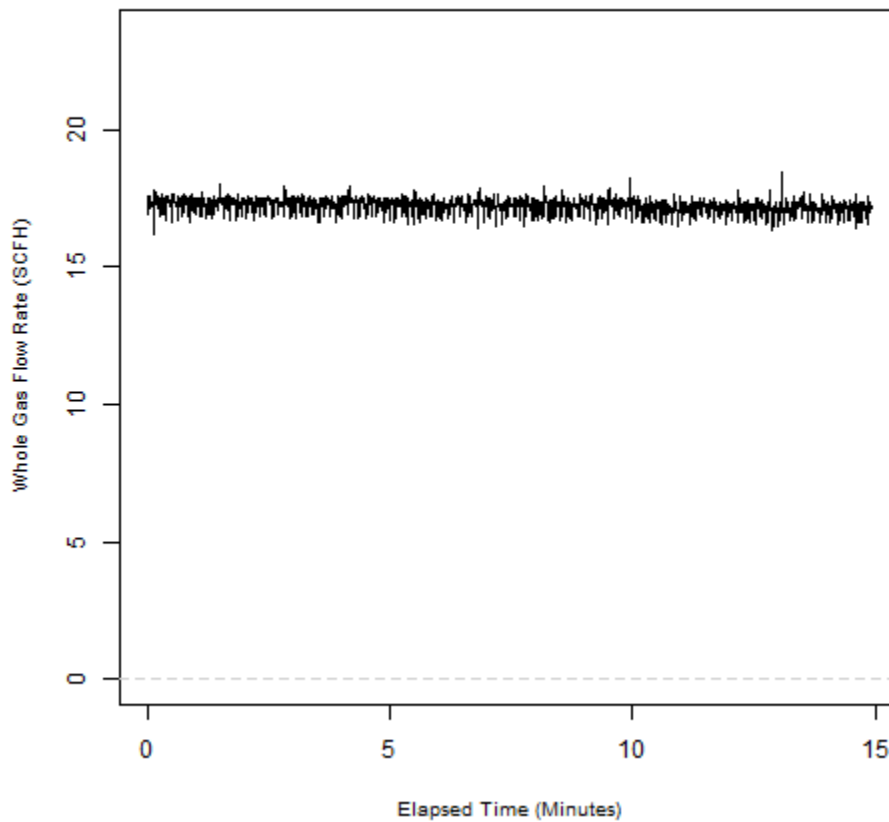
Avg. Emission rate (whole gas):	17.4 scf/h
Avg. Emission rate (methane):	15.3 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Oil Level Control
Manufacturer/Model (blinded):	G01
Supply pressure:	22 psig
Assessment of controller operation:	equipment issues



RB01-PC13 [38] Region: GC

Device characteristics:

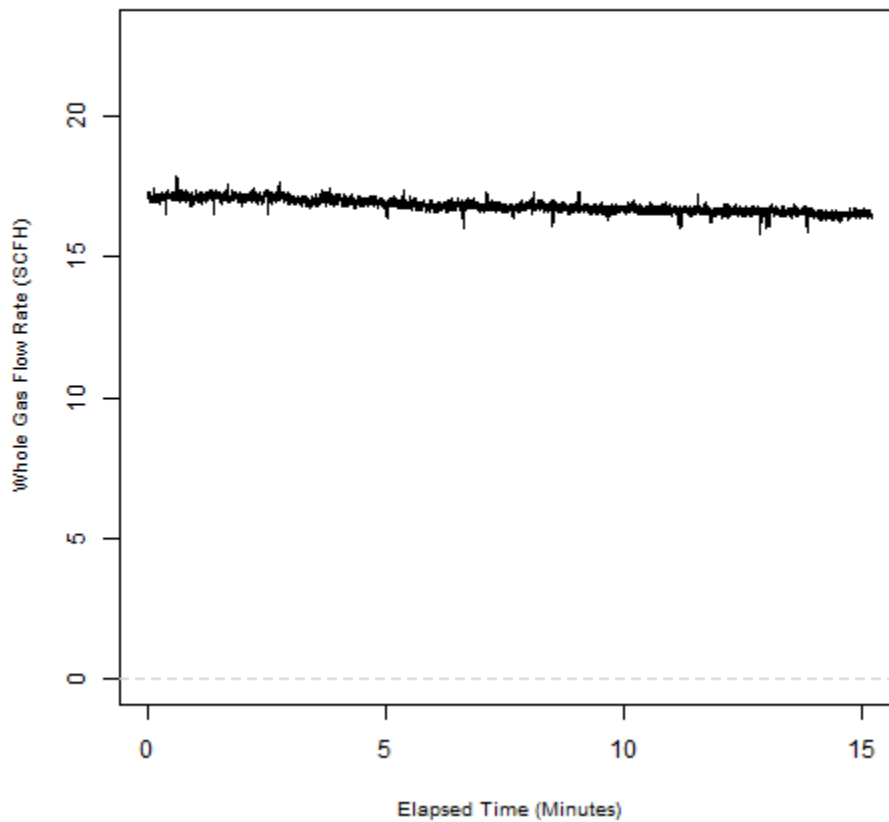
Avg. Emission rate (whole gas):	17.2 scf/h
Avg. Emission rate (methane):	14.2 scf/h
Controller application:	Level
Basic Application:	Dehydration System
Detailed Application:	Contact Tower- Liquid Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	25 psig
Assessment of controller operation:	equipment issues



VF01-PC22 [39] Region: AP

Device characteristics:

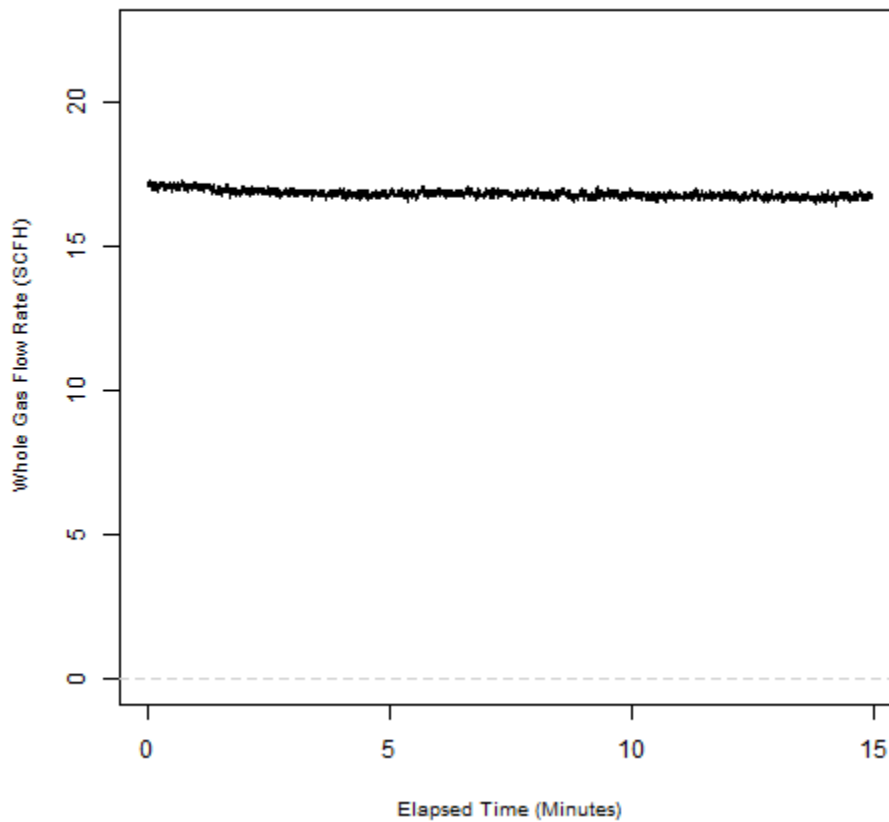
Avg. Emission rate (whole gas):	16.8 scf/h
Avg. Emission rate (methane):	16.4 scf/h
Controller application:	Temperature
Basic Application:	Process Heater
Detailed Application:	Gas Flow to Process Heater
Manufacturer/Model (blinded):	I04
Supply pressure:	7 psig
Assessment of controller operation:	equipment issues



RB01-PC34 [40] Region: GC

Device characteristics:

Avg. Emission rate (whole gas):	16.8 scf/h
Avg. Emission rate (methane):	13.8 scf/h
Controller application:	Level
Basic Application:	Separator
Detailed Application:	Separator - Water Level Control
Manufacturer/Model (blinded):	M01
Supply pressure:	32 psig
Assessment of controller operation:	equipment issues



B Methane Emissions from Liquids Unloadings

B.1 Methods for site selection

Goals and Overall Sampling Strategy. Sampling of emissions from gas well liquids unloadings was conducted in four major regions (Appalachian, Gulf Coast, Mid-continent, Rocky Mountain). The regions are shown in Figure B-1.

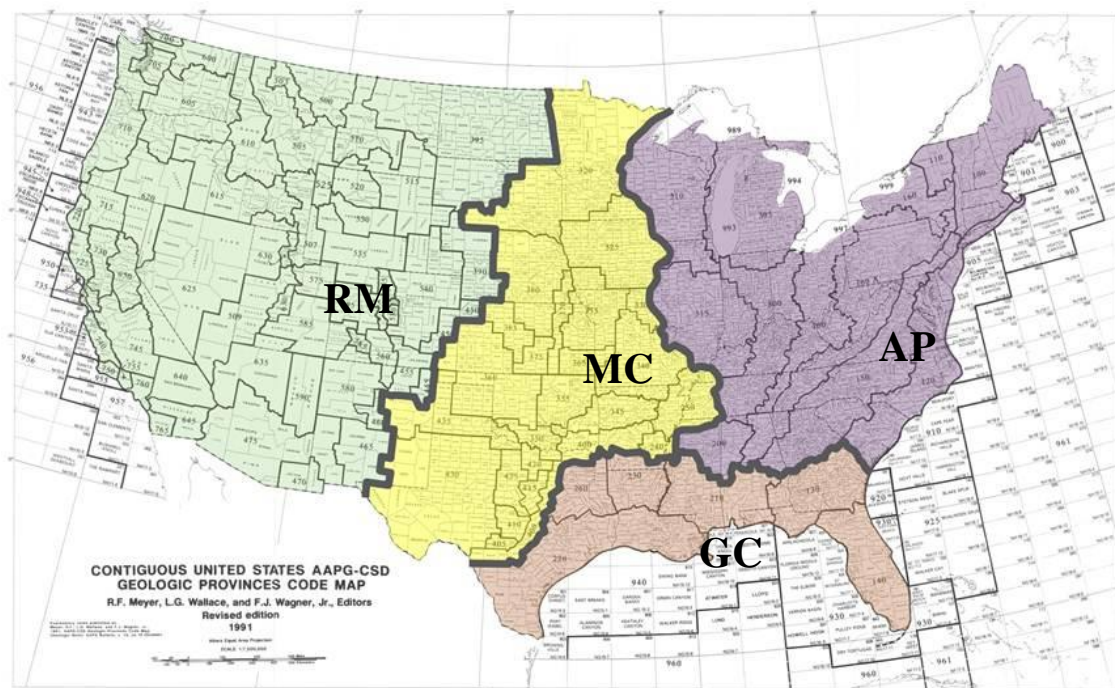


Figure B-1. Basins of the American Association of Petroleum Geologists (AAPG) divided into 4 Major Regions for this study: AP=Appalachian; GC=Gulf Coast; MC=Mid-Continent; RM=Rocky Mountain.

It was anticipated that in each of the four regions, gas wells with and without plunger lifts would be sampled, and that within each of these categories, there would be a range of unloading frequencies, durations and liquid production rates. To adequately sample regions, well types (plunger and without plunger) and unloading event characteristics, it was anticipated that measurements of unloading emissions from approximately 100 different wells would be required.

Selection of Site Visit Duration and Scope. With a goal of 100 well unloading measurements, the project team conducted approximately 20 one-week visits to natural gas production regions with unloading emissions. It was anticipated that 5 wells could be sampled in a typical week. Production basins with the highest emissions, as reported through the U.S. EPA's Greenhouse Gas Reporting Program, were targeted. Each week of sampling was conducted with a single company in a single basin location.

Selection of Basins. Basins in which sampling was conducted were selected based on emissions reported through the EPA's Greenhouse Gas Reporting Program (GHGRP, Reporting Year 2012) Figure B-2 shows Basins reporting unloading emissions through the GHGRP. Any basin colored blue had reported unloading emissions; uncolored (white) basins had no reported emissions. The darkest blue indicates basin total emissions for all reporters in excess of one million metric ton of CO₂e annually (based on a Global Warming Potential for methane of 21), medium blue indicates basin total emissions between 100,000 and 1,000,000 metric tons (MT) CO₂e annually, and lightest blue are basins that reported less than 100,000 MT CO₂e annually. There were 27 basins that reported unloading emissions in 2012.

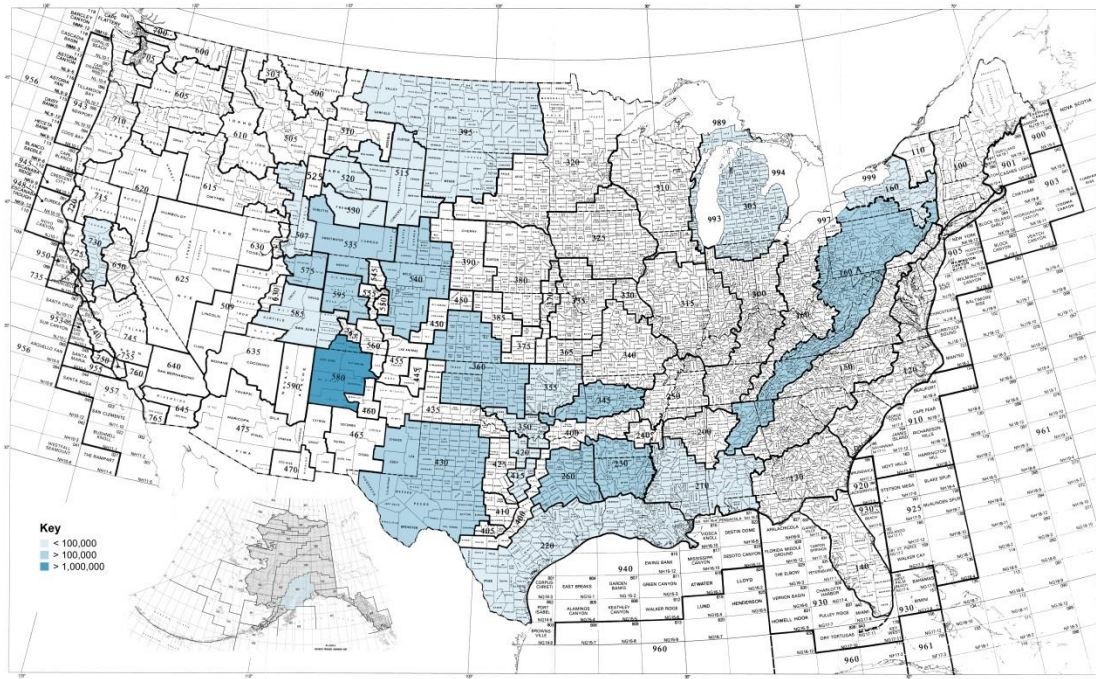


Figure B-2. Basins of the American Association of Petroleum Geologists (AAPG) where unloadings were reported in 2012 to the EPA GHGRP.

Ten companies participating in this work reported 60% of the total unloading emissions for GHGRP reporting year 2012, and account for 26% of the wells that reported emissions.

Table B-1. Spatial distribution of total unloading emissions reported in the GHGRP compared to the spatial distribution of emissions reported by companies that provided sampling sites

Region	AAPG Basin	Total GHGRP Emissions for 2012 (MT CO ₂ e)	% of total Basin emissions accounted for by companies providing sampling sites	% of total Basin wells that have unloadings accounted for by companies providing sampling sites
Appalachian	160A - Appalachian Eastern Overthrust	413,623	14.9%	14.5%
	220 - Gulf Coast Basin	74,525	31.6%	40.1%
GC	230 - Arkla Basin	148,126	8.4%	36.9%
	260 - East Texas Basin	242,828	82.4%	65.5%
	345 - Arkoma Basin	477,471	20.0%	26.6%
MC	350 - South Oklahoma Folded Belt	972	6.2%	6.9%
	360 - Anadarko Basin	310,355	1.2%	13.2%
	415 - Strawn Basin	43,050	48.3%	54.9%
	420 - Fort Worth Syncline	32,933	14.4%	39.1%
	430 - Permian Basin	179,707	1.8%	3.5%
RM	507 - Central Western Overthrust	42,505	14.0%	32.4%
	530 - Wind River Basin	4,743	84.5%	23.4%
	535 - Green River Basin	182,427	24.0%	40.2%
	540 - Denver Basin	102,335	6.1%	22.3%
	575 - Uintah Basin	149,584	8.9%	25.6%
	580 - San Juan Basin	2,315,772	96.4%	87.3%
	595 - Piceance Basin	943,554	79.2%	12.1%
Total US		5,846,634	59.5%	25.5%

Selection of Company and Basin Locations The Study Team, consisting of URS and University of Texas personnel, was solely responsible for the selection of regions and Basins in which to sample. For most basins, more than one of the ten participant companies has reported unloading emissions. If every participant company were visited in each basin where any participant unloading emissions were reported, there would have been 52 weeks of site visits. Since project scope and budget called for approximately 20 sample weeks, a subset of all possible participant sites were selected for sampling.

The selection of company sites required a balance among a number of goals. One goal was to sample at least 3 companies in each major region (AP, GC, MC, RM) shown in Figure B-1. A second goal was to sample the basins with the largest reported emissions in the GHGRP. A third goal was to be able to sample each of the ten participant companies at least once. All companies that reported wells with unloading emissions were sampled in this program.

Once a basin was targeted for sampling, selection of the particular company to visit started with the participant company with the largest reported emissions in the basin, unless that company had already been sampled elsewhere, or unless one of the other participant companies only reported emissions in that single basin. Figure B-3 shows the basins that were sampled during this measurement effort in late 2013 and early 2014. All basins with emissions of more than 100,000 MT CO₂e/yr from unloadings, as reported through the GHGRP, were sampled, with the exception of basin 360. The one company targeted for basin 360 did not have had any unloading events for the week selected for a site visit to Basin 360.

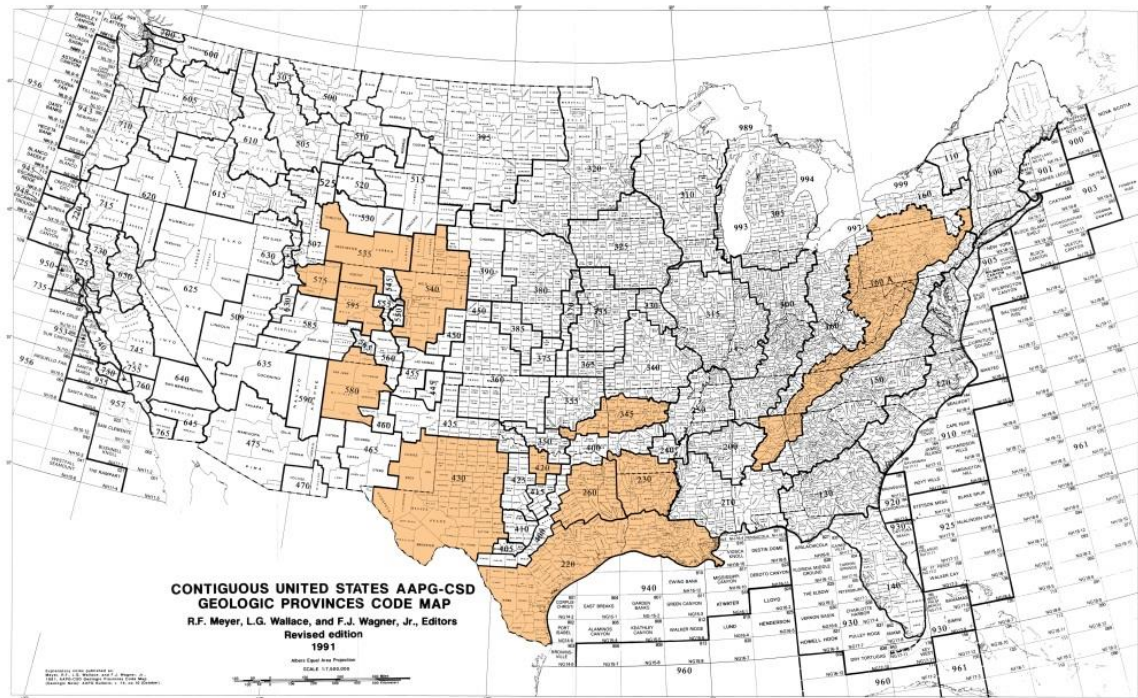


Figure B-3. Basins of the American Association of Petroleum Geologists (AAPG) where unloadings measurements were made in this work.

Once a Basin and company to be sampled was selected, local contacts for participant companies provided descriptions of the types of unloadings and typical frequencies expected. No companies refused a site visit. Once at a site, the Study Team measured emissions from as many wells as could be visited and measured in the week. In some cases this involved sampling every unloading that occurred during the week for the company being visited. When more unloadings were available than could be sampled during a week, the Study Team selected which wells to visit.

B.2 Corrections to Instrument Flow Measurements based on Temporary Stack Size and Gas Composition

When safe and technically possible, the flow measurements of gas volumes released during liquid unloadings were taken using a temporary stack affixed to a tank vent that

was equipped with a gas velocity measurement instrument (Fox Thermal Instruments, Model #FT3). On sites where the unloading flow was directed to an open top blowdown tank, rather than a fixed roof tank, a length of pipe was inserted into the piping to the open top tank that was used for the unloading, allowing for measurement of the flow into the tank. The Fox #FT3 device measured velocity over a 1.4” measurement length (with 0.5 inch thickness), which was centered in the temporary stack or pipe. The average-center line velocity measured by the Fox #FT3 was converted to an average velocity in the temporary stack or pipe by assuming that the velocity distribution was well-represented by a one-seventh power law velocity distribution.

$$\frac{v(r)}{v_{max}} = \left(1 - \frac{r}{R}\right)^{1/7} \quad (\text{B2.1})$$

where $v(r)$ is the velocity at distance r from the tube centerline, and R is the radius of the temporary stack or pipe. Typically, the power law order (in this case $1/7$) is a function of the Reynolds number of the flow. However, over a wide range of Reynolds numbers, the differences in the predicted stack velocities are not sensitive to the assumed order of the power law function (between $1/7$ and $1/9$) except near the stack or pipe wall. In this work, the measurements were made near the center-line, and the effect near the stack or pipe wall would be expected to be minimal.

The specific relationship between the measured velocity and the overall average velocity depended on the size of the stack or pipe, since the 1.4 inch Fox #FT3 probe measured a different proportion of the cross sectional flow for different sized pipes. For all stacks and pipes, the ratio of the center-line (maximum) velocity to the average velocity over the entire pipe was 0.82. So, if the probe had only measured centerline velocity, the ratio of the average velocity to measured centerline velocity would be 0.82. The actual ratio of the measured velocity to the maximum velocity was dependent on the fraction of the diameter of the pipe that was sampled by the 1.4 in. probe. The correction factor was calculated using the one seventh-order power law distribution, and an assumption that the

probe was exactly centered. The parameters for the pipes and stacks used in this study are summarized in Table B-2.

Table B-2. Correction factors used to account for the difference in the measured velocity and the average velocity through the pipe or stack. Note that the 2.5” nominal diameter stack was only used for the calibration of some of the Fox #FT3 devices and was not used for in-field measurements during the study.

Nominal Stack Diameter (in)	Stack Internal Diameter (in)	Proportion of Stack Diameter Measured (r/R)	Ratio of Measured Velocity to Center-Line Velocity (Average velocity = 0.82 * centerline velocity)
2	2.060	0.68	0.93
2.5	2.469	0.57	0.95
3	3.068	0.46	0.96
6	5.76	0.24	0.98
8	7.90	0.18	0.98

The volumetric flow rate is the average velocity in the stack or pipe multiplied by the cross-sectional area available for flow. In the measurements made in this study, the cross-sectional area available for flow is the cross-sectional area of the stack or pipe minus the area obstructed by the flow probe, as shown in Figure B-4. The area available for flow through each of the stack sizes used in this study is shown in Table B-3.



Figure B-4. Fox #FT3 velocity probe centered in a 2” nominal diameter stack.

Table B-3. Parameters for stacks and pipes used in this study and the unobstructed cross-sectional area for flow, accounting for the Fox #FT3 probe insertion. Note that the 2.5” nominal diameter stack was only used for the calibration of some of the Fox #FT3 devices and was not used for in-field measurements during the study.

Nominal Stack Diameter (in)	Stack Area (in ²)	Area Obstructed by Probe (in ²)	Unobstructed Area (in ²)
2	3.33	0.57	2.76
2.5	4.79	0.67	4.11
3	7.39	0.82	6.57
6	26.07	1.50	24.6
8	49.0	2.03	47.0

Thus, the instrument reported flow in standard cubic feet per hour (scf/h) was calculated based on the instrument reported velocity and the cross-sectional area available for flow through the stack or pipe:

$$scf/h_{inst} = v_{inst} * \frac{0.82}{v_m/v_{max}} * \frac{A_c}{144} \quad (B2.2)$$

where v_{inst} is the instrument reported velocity in feet per hour, v_m/v_{max} is the ratio of measured velocity to center-line velocity (Table B-2), and A_c is the unobstructed cross-sectional area of the stack or pipe in square inches (Table B-3). The overall factor converting instrument reading to volumetric flow for each stack and pipe size is summarized in Table B-4.

Table B-4. Overall conversion factor for velocity measurement (ft/h) to flow rate (scf/h) for each stack and pipe size used in the project. Note that the 2.5” nominal diameter stack was only used for the calibration of some of the Fox #FT3 devices and was not used for in-field measurements during the study.

Nominal Stack Diameter (in)	Factor (B)
2	0.0169
2.5	0.0246
3	0.0390
6	0.143
8	0.273

The Fox FT#3 flow measurement devices were calibrated on pipes with sizes that were not necessarily the same as those utilized in the field measurements for unloadings. The probes were calibrated by the manufacturer by sending a known volume of methane through a pipe of known diameter and cross sectional area. For this study, calibrations were made on different Fox #FT3 instruments using the pipes specified as 2.5” and 3” nominal diameter in Tables B-2 and B-3. Thus, the flow rate calculated by the instrument (using a pipe which was typically a different diameter than the pipe used for calibration) had to be scaled by the ratio of the factor (Table B-4) for the field pipe to the calibration pipe.

In addition, the Fox #FT3 instruments were calibrated by the manufacturer (Fox Thermal Instruments) using pure methane; instrument flow rates were corrected to account for the measured gas compositions, which varied from site to site. Because the flow meter measurement is based on thermal conductivity, the composition correction was based on the relative thermal conductivities of the gas used by pneumatic devices at each site and the pure methane used as a calibration gas.

$$scf/h = scf/h_{inst} \left(\frac{k_{CH_4}}{k_{gas}} \right) \left(\frac{B_{field}}{B_{cal}} \right) \quad (B2.3)$$

where scf/h_{inst} is the raw instrument flow rate reading in standard cubic feet per hour, k_{CH4} is the pure component thermal conductivity (W/m*K) for methane at standard conditions (70°F and 14.7 psia), and k_{gas} is the thermal conductivity of the gas sampled at standard conditions. The B factors are the values in Table B-4, which convert the measured velocity to a flow rate. The thermal conductivity of the sampled gas (k_{gas}) was calculated as a molar weighted average:

$$k_{gas} = \sum_{i=1}^7 k_i n_i \quad (B2.4)$$

where n_i is the mole fraction of species i in the gas sample for the site and k_i is the pure component thermal conductivity of the species (<http://webbook.nist.gov/chemistry/fluid/>) under standard conditions. For this work, the pure component species considered were methane, ethane, propane, nitrogen, air, and carbon dioxide. All higher hydrocarbons with a carbon count of four or greater were lumped with butane for purposes of the gas composition correction.

Table B-5. Thermal conductivity of measured species at standard conditions (14.7 psia and 70°F).

Species	Thermal Conductivity (W/m*K)
Methane	0.033759
Ethane	0.020491
Propane	0.017884
Butane +	0.016181
Nitrogen	0.025473
Carbon Dioxide	0.016331

B.3 Unloading emission data

Table B-6. Unloading emissions from wells with automatically triggered plunger lifts.

Well Characteristics					Emissions Data				
Well	Region	V or H*	Gas prod. rate (scf/d)	CH ₄ in gas (%)	CH ₄ emitted per event (scf)	Events sampled	Avg. event duration (s)	Events per year for well reported by operator	Emiss. per year for well based on events reported by operator (MMscf CH ₄)
UBB 42 0101	RM	V	170,000	75.7	914	6	147	1982	1810
UBB 42 0201	RM	V	100,000	78.7	8,621	2	1208	1069	9220
UBB 42 0401	RM	V	200,000	81.9	64	7	389	2546	163
UBB 42 0501	RM	V	140,000	82.9	659	18	130	606	399
UBB 42 0601	RM	V	140,000	83.4	7,278	5	1208	2686	19500
UBB 42 0701	RM	V	170,000	81.1	103	9	276	184	19
UBB 42 0801	RM	V	155,000	80.5	1,695	3	433	2048	3470
UBB 42 0901	RM	V	150,000	91.5	91	6	70	964	88
UBB 42 1001	RM	V	162,000	91.5	209	5	36	715	149
UBB 42 1101	RM	V	175,000	80.0	1,534	76	262	1011	1550
UBB 42 1201	RM	V	90,000	80.4	611	7	617	573	350
UBB 43 0101	RM	V	180,000	78.5	1,296	8	177	2873	3720
UBB 43 0301	RM	V	190,000	81.7	64	6	137	4698	301
UEY 41-0101	MC	V	38,000	97.8	312	4	400	6570	2050
UEY 41-0201	MC	V	90,000	97.8	129	2	274	2389	308
UEY 41-0301	MC	V	129,000	97.8	215	4	191	7509	1614
UEY 41-0401	MC	V	107,000	97.8	549	1	206	3893	2137
UBB 50 2601	RM	V	54,000	76.8	915	2	317	643	588
UBB 50 2701	RM	V	47,000	77.7	1,998	1	849	4252	8495
UBB 50 2801	RM	V	32,000	78.2	56	23	33	4051	227
UBB 50 2901	RM	V	15,000	83.0	993	14	90	2482	2464
UBB 50 3001	RM	V	26,000	81.9	237	7	692	650	154
UBB 50 3101	RM	V	46,000	80.6	58	17	73	1963	114
UBB 50 3201	RM	V	46,000	76.6	1,325	1	309	528	700
UEF-49-0501	GC	V	18,000	82.4	428	2	123	4238	1810

* (V)ertical or (H)orizontal.

Table B-7. Unloading emissions from wells with manually triggered plunger lifts

Well Characteristics			Emissions Data						
Well	Reg.	V or H	Gas prod. rate (scf/d)	CH ₄ in gas (%)	CH ₄ emit. per event (scf)	Events samp.	Avg. event duration (s)	Events per year for well reported by operator	Emiss. per year for well based on events reported by operator (MMscf CH ₄)
UBB-45-0101	RM	V	20,000	77.4	9,674	1	727	30	290
UBB-45-0201	RM	V	20,000	77.4	11,678	1	1,529	50	584
UBB-45-0202	RM	V	35,000	77.4	11,783	1	1,303	50	589
UBB-45-0203	RM	V	10,000	77.4	4,703	1	1,522	50	235
UBB-45-0204	RM	V	25,000	77.4	3,641	1	1,708	50	182
UBB-45-0301	RM	V	16,000	77.4	5,612	1	1,416	2	11
UBB-45-0302	RM	V	45,000	77.4	4,008	1	1,219	2	8
UBB-45-0401	RM	V	25,000	77.4	16,852	1	3,714	2	34
UBB-45-0501	RM	V	13,000	77.4	8,057	1	3,847	5	40
UBB-47-0301	GC	V	80,000	86.5	3,937	1	7,672	2	8
UDN-44-0203	RM	V	125,000	88.8	4,737	1	443	3	14
UDN-44-0304	RM	V	43,000	90.7	14,069	1	540	6	84
UDN-44-0405	RM	V	343,000	89.1	8,289	1	417	3	25
UDN-44-0506	RM	V	231,000	89.9	6,459	1	528	1	6
UDN-44-0507	RM	V	248,000	88.4	8,639	1	588	*	
UEF-02-0201	MC	V	41,000	81.1	8,290	1	2,271	13	108
UEF-02-0202	MC	V	15,000	84.3	6,083	1	5,637	10	61
UEF-02-0203	MC	V	24,000	82.5	11,958	1	2,864	12	143
UEF-02-0204	MC	V	35,000	78.4	6,272	1	1,780	8	50
UEF-02-0205	MC	V	24,000	81.9	14,570	1	3,458	8	117
UEF-02-0206	MC	V	38,000	85.6	21,255	1	4,996	4	85
UJR-46-0101	RM	V	86,000	86.4	1,665	1	1,011	5	8
UJR-46-0401	RM	V	36,000	86.4	993	1	231	6	6
UJR-46-0501	RM	V	37,000	86.4	6,744	1	560	16	108
UJR-46-0601	RM	V	30,000	86.4	1,220	1	198	1	1
UJR-46-0701	RM	V	34,000	86.4	1,261	1	339	11	14
UJR-46-0801	RM	V	51,000	86.4	22,364	1	3,926	3	67
UJR-46-1001	RM	V	110,000	86.4	8,101	1	3,149	2	16
UJR-46-1101	RM	V	118,000	86.4	2,663	1	559	15	40
UJR-46-1201	RM	V	19,000	86.4	21,060	1	1,289	8	168
UMB-06-0101	MC	V	23,000	83	4,805	1	790	2	10

Table B-7 (continued).

Well Characteristics			Emissions Data						
Well	Reg.	V or H	Gas prod. rate (scf/d)	CH ₄ in gas (%)	CH ₄ emit. per event (scf)	Events samp.	Avg. event duration (s)	Events per year for well reported by operator	Emiss. per year for well based on events reported by operator (MMscf CH ₄)
UMB-06-0401	MC	V	226,000	81.5	403	1	169	1	0.5
UMB-06-0501	MC	V	4,000	82.1	1,311	1	474	2	3
UMB-06-0601	MC	V	30,000	82.5	1,245	1	387	10	12
UMB-06-0901	MC	V	38,000	77.5	1,862	1	688	10	19
USH-42-0301	AP	H	900,000	97.3	223	1	86	7	2
USH-45-0101	RM	V	160,000	87.8	18,277	1	1,314	2	37
USH-45-0103	RM	V	168,000	87.8	47,119	1	1,181	3	141
USH-45-0105	RM	V	87,000	87.8	8,924	2	2,720	15	134
USH-45-0201	RM	V	119,000	87.8	26,668	1	1,900	1	27
USH-45-0202	RM	V	166,000	87.8	49,273	1	1,447	11	542
USH-45-0203	RM	V	151,000	87.8	15,834	2	1,638	39	618
UTG-44-0201	AP	V	6,000	95.7	4,313	1	10,214	20	86
UTG-44-0301	AP	V	10,000	95.7	8,622	1	2,609	24	207
UTG-44-0401	AP	V	4,000	95.7	4,398	1	3,213	24	106
UTG-44-0501	AP	V	10,000	95.7	3,534	1	1,240	24	85
UTG-44-0601	AP	V	17,000	95.7	3,964	1	1,060	52	206
UTG-44-0701	AP	V	6,000	95.7	10,542	1	9,807	12	127

Table B-8. Unloading emissions from wells without plunger lifts

Well Characteristics			Emissions Data						
Well	Reg.	V or H	Gas prod. rate (scf/d)	CH ₄ in gas (%)	CH ₄ emitted per event (scf)	Events samp.	Avg. event duration (s)	Events per year for well reported by operator	Emiss. per year for well based on events reported by operator (MMscf CH ₄)
UBB-47-0101	GC	V	150,000	92.7	555	2	1,518	2	1
UBB-47-0201	GC	V	100,000	80.7	6706	2	3,417	4	27
UBB-47-0401	GC	V	100,000	86.5	2745	1	2,104	1	3
UCG-03-0101	GC	V	162,078	96.1	12237	1	6,762	48	587
UCG-03-0102	GC	V	155,279	96.1	13761	1	6,016	6	83
UCG-03-0103	GC	V	160,791	96.1	24085	1	7,919	12	289
UCG-03-0201	GC	V	153,467	93.3	16029	1	3,952	3	48
UCG-03-0202	GC	V	269,018	93.3	24544	1	4,739	3	74
UCG-03-0203	GC	V	167,000	96.1	16056	1	2,504	185	1975
UCG-03-0204	GC	V	43,748	93.3	9942	1	5,120	81	805
UCG-03-0301	GC	V	102,050	93.3	21342	1	9,819	27	576
UCG-03-0302	GC	V	151,000	96.1	11436	1	4,308	151	1269
UCG-03-0401	GC	V	123,017	96.1	10696	1	1,270	9	96
UCG-03-0402	GC	V	67,718	96.1	16487	1	2,662	45	742
UEY-41-0601	MC	H	400,000	95.2	73417	1	11,782	1	73
UMB-06-0701	MC	V	25,000	74.3	3509	1	1,841	5	18
UMB-06-0801	MC	V	20,000	72.8	6460	1	3,580	3	19
UMB-06-1101	MC	V	21,000	81.3	6083	1	1,021	12	73
UMB-06-1201	MC	V	4,787	78.4	1951	1	964	1	2

Table B-8 (continued).

Well Characteristics			Emissions Data						
Well	Reg.	V or H	Gas prod. rate (scf/d)	CH ₄ in gas (%)	CH ₄ emitted per event (scf)	Events samp.	Avg. event duration (s)	Events per year for well reported by operator	Emiss. per year for well based on events reported by operator (MMscf CH ₄)
USH-42-0501	AP	H	230,000	97.3	1423	1	1,252	6	9
USH-45-0102	RM	V	93,000	87.8	9409	1	2,411	19	179
USH-45-0104	RM	V	86,000	87.8	20967	1	805	5	105
USH-47-0101	MC	H	365,200	95.3	41919	1	10,009	4	168
USH-47-0201	MC	H	224,404	96.7	75974	2	14,525	95	7218
USH-47-0301	MC	H	147,231	96.7	57793	1	16,161	101	5837
USH-47-0401	MC	H	182,770	97.5	27055	1	7,766	84	2273
USH-47-0601	MC	H	258,500	94.8	47037	1	3,971	70	3293
USH-47-0701	MC	H	246,356	97.8	13483	1	9,741	45	6068
USH-47-0801	MC	H	88,217	97.9	97937	1	5,530	2	196
UTG-44-0101	AP	V	10,000	95.7	4007	1	629	12	48

*Of the two events measured for this well, one was done without the well being shut-in before the event; since liquids removal was low during this unloading, a second unloading, was done 3 days later with the well shut-in for the 3 days between events. According to the well operator this shut in is required for approximately 6 of the 95 events during a year. The average emissions is therefore a weighted average of the first event (typical of 89 of the 95 events per year) and the much larger event (6 of 95 events per year).

B.4 Statistical analyses of variability in unloading emission measurements

The emission measurement data were combined with well characteristics reported by the host companies to identify possible explanatory variables for the frequency of unloading events and annual unloading emission totals. A natural logarithm (ln) transform was applied to the total gas annual emission values owing to the skewness in this variable. The logarithm transform maintains the ordering of observations but reduces the influence of the larger values on the calculated statistics. The statistical Pearson linear correlations between annual emissions and number of events per year with several other statistical variables were calculated. The correlation is a number between -1.0 and 1.0 that is a measure of the linear association between two variables. A positive correlation between two variables suggests that generally if one observation of the first variable is higher than the average for that variable, then the corresponding value the second variable is also higher than average. A negative correlation between two variables suggests that generally if the first variable is higher than the average for that variable, then the corresponding value of the second variable is lower than average. One must use care in concluding that there is a causal relationship underlying a high positive or negative correlation. One confounding factor could be that there are outliers in the data that have disproportionate impact on average values and therefore in producing a calculated correlation. The natural logarithm transform used on the annual emission total helps to address this. Care must also be used in drawing a conclusion about a correlation close to zero, as a strong relation could exist between variables with correlation = 0.0 that is nonlinear.

Associated with a calculated linear correlation is a probability value (p-value) that represents the approximate probability that a correlation as large as the one calculated could have been the result of a random set of data with no underlying correlation. An individual correlation result with a p-value of 0.05 or less means that the probability that a value as large as the calculated correlation would result if the variables had a random association is small, and thus we conclude the association is likely not random.

The correlations were calculated between company reported *Events per Year*, measured *Event Duration*, estimated *Whole Gas SCF per Year*, and the natural logarithm of *Whole Gas SCF per Year* with 7 well characteristic variables. The results are shown in Table B-9. The correlations with p-values below 0.05 are shown in Table B-10. Although several pairings of emissions variables and well variables appear to be statistically significant, the scatter plot graphs and linear fits suggest most of these relationships do not explain more than a few percent in the variability of emissions, which is the correlation-squared or R^2 in linear regression. Well *Total Depth* is an exception, as shown in the line fit graph in Figure B-5. *Total Depth* explains 14% ($R^2 = 0.14$) of the variability in $\ln(\text{Whole Gas SCF per Year})$ and there is a visible downward trend in the data suggesting emissions decrease with well depth. This may or may not be related to a tendency in deeper wells being newer and thus less prone to unloadings. The variable for well *Age* was statistically significantly positively correlated with the estimated number of *Events per Year* and not-statistically significant but still positively correlated with *Whole Gas SCF per Year*.

Table B-9. List of study measurements and estimates compared with reported well characteristics

Study measurements	Well characteristics
Event duration	Surface flow line pressure
Events per year	Static Shut-in pressure
Whole Gas SCF per Year	Total Depth
Ln(Whole Gas SCF per Year)	Production SCF per day
	Volume (depth * diameter * diameter / 4)
	Age of well

Table B-10. Pairwise linear correlations

Y Variable	X Variable (study measurements in bold)	# of paired obs.	Linear Correlation	R ²	p-values
Event duration	CH4 scf / event	106	0.6368	40.60%	0.0001
Event duration	CH4 pct	106	0.4434	19.70%	0.0001
Event duration	age	81	-0.1954	3.80%	0.0805
Events per year	Total Depth	105	-0.3926	15.40%	0.0001
Events per year	Event duration	106	-0.2794	7.80%	0.0037
Events per year	volume	105	-0.2438	5.90%	0.0122
Events per year	CH4 scf / event	106	-0.2244	5.00%	0.0208
Events per year	Static Shut in pressure	86	-0.2413	5.80%	0.0252
Events per year	Surface flow line pressure	96	-0.1881	3.50%	0.0665
ln(Whole gas scf/yr)	Events per year	106	0.4463	19.90%	0.0001
ln(Whole gas scf/yr)	volume	105	-0.3138	9.80%	0.0011
ln(Whole gas scf/yr)	CH4 scf / event	106	0.2881	8.30%	0.0028
ln(Whole gas scf/yr)	Event duration	106	0.2209	4.90%	0.0229
Whole gas scf / year	Events per year	106	0.3303	10.90%	0.0005
Whole gas scf / year	Total Depth	105	-0.2616	6.80%	0.007
Whole gas scf / year	age	81	0.2508	6.30%	0.0239
Whole gas scf / year	volume	105	-0.177	3.10%	0.0708
Whole gas scf / year	CH4 scf / event	106	0.1759	3.10%	0.0713

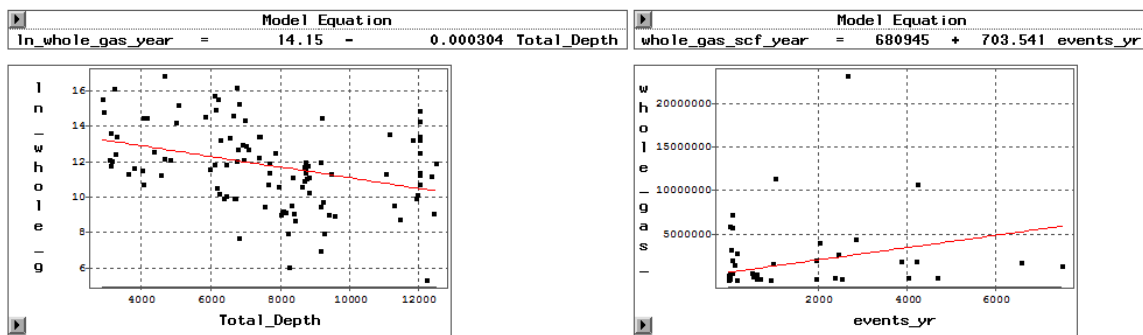


Figure B-5. Examples of linear fits for two significantly related variables (*left*: ln whole gas emissions vs. well depth for all wells; *right*: ln whole gas emissions vs. frequency of unloading for automated plunger lift wells).

Additional statistical analyses were done, comparing the observed emissions per event, to emissions per event that would be predicted based on EPA emission estimation methods described in Technical Support Documents available for GHGRP reporting, (http://www.epa.gov/ghgreporting/documents/pdf/2010/Subpart-W_TSD.pdf). Briefly, these emission estimation methods assume that an unloading event vents, at a minimum, the entire volume of the well bore and that the well bore is entirely filled with gas at the shut-in pressure. If, for a plunger lift well, the event lasts more than 30 minutes, the vent rate for the period after 30 minutes is assumed to be at the production rate. If, for a non-plunger lift well, the event lasts more than 60 minutes, the vent rate for the period after 60 minutes is assumed to be at the production rate. The comparisons with the emission estimation method were done for wells on which all of the data required for the emission estimation method were available.

For wells without plunger lift (18 wells with all data available), the observations were higher (statistically significant difference) than the estimates, however, linear correlations, shown in Figure B-6, led to a slope that was not statistically significant. If the two highest observed emissions events are removed from the analysis (events circled in Figure B-6), the correlation does become statistically significant.

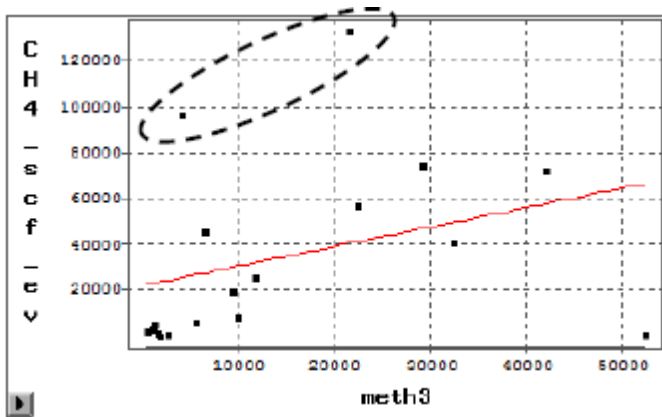


Figure B-6.- Comparison of emission predictions using EPA emission estimation methods to observed emissions per event for wells without plunger lift.

For wells with plunger lifts (66 wells), the emission estimates average 20,200 scf/event as compared to an average of 7,100 scf/event for the measurements. The difference is statistically significant ($p=0.0001$), however the predicts and the observations are positively correlated. Figure B-7 shows the comparison.

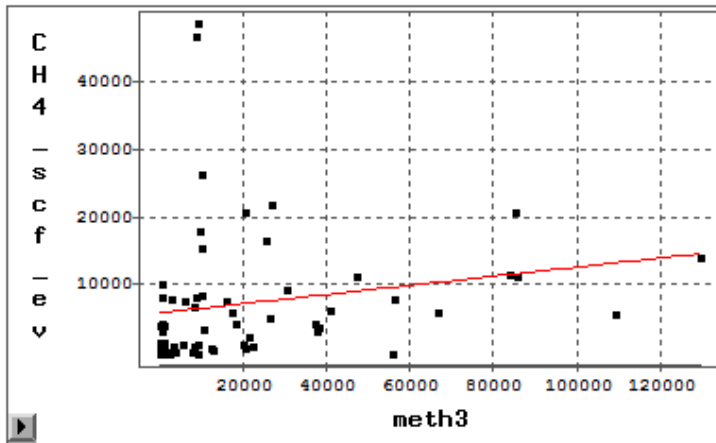


Figure B-7. Comparison of emission predictions using EPA emission estimation methods to observed emissions per event for wells with plunger lift.

B.5 Estimates of emissions from gas well liquid unloadings in the United States

Emission measurements from a limited set of samples can be used to estimate national emissions by multiplying the average emission measurement by the number of times that emission occurs on the national scale. Often the emission measurement is referred to as an “emission factor” or EF, and the data used to scale up the emissions is called the activity factor (AF). Emissions are calculated as:

$$EF_i * AF_i = ER_i \tag{B5.1}$$

where:

EF_i = Emission Factor for region i

AF_i = Activity Factor for region i

ER_i = resulting Emission Rate total for region i

For this work, the activity factors are national counts of natural gas well unloading events. The activity data and the emission factor data are stratified at two levels. First, wells are classified as either with or without plunger lift. Then, within each well category (with or without plunger lift) wells are grouped based on the frequency of unloadings (events per year). The emission factor strata for which activity data are needed are shown in Table B-11.

Table B-11. Categories of wells for which emissions were measured

Well type (Annual Frequency of events)	Measured EF, scf methane per event (95% confidence range)
Plunger Wells (events<100)	9,450 (6,900-12,400)
Plunger Wells (events≥100)	1,200 (500-2,100)
Non Plunger Wells (events<10)	21,500 (9,600-37,800)
Non Plunger Wells (10≤events<50)	24,100 (8,700-50,400)
Non Plunger Wells (50≤events<200)	35,000 (18,700-53,000)
Non Plunger Wells (events≥200)	Not measured, assume 35,000 (see main text) (18,700-53,00)

Activity Data: The primary source of activity data used in this work is a survey of unloading event count collected from companies that participated in this work. These survey data will be supplemented by data on well counts from the GHG Mandatory Reporting Program, for reporting year 2012, released in 2013 (referred to here as EPA 2012 GHGRP) and the 2012 GHG National Emission Inventory, released in 2014 (referred to here as the 2012 GHG NEI).

The EPA 2012 GHG NEI reports that 60,810 natural gas wells have liquids unloadings, out of an estimated 470,913 natural gas wells in the United States (not including oil wells with associated gas production). This represents 13% of gas wells in the EPA 2012/2014 GHG NEI. Of these 60,810 wells, 23,503 are reported as having plunger lifts and 37,307 are reported as not having plunger lifts.

More granular and detailed data is available as part of the EPA 2012 GHGRP. All operators of U.S. upstream petroleum and natural gas production are required to report under the GHGRP, so long as their total emissions from all sources exceeds 25,000 MT CO₂e/yr for an entire basin. EPA estimated in the original publication of the Rule that more than 85% of all GHG emitters would have to report, though the percentage may be higher in the oil and gas production sector, given that reporting facilities are defined as large basins. The latest data available from the GHG Mandatory Reporting rule is published in the EPA's Facility Level Information on GreenHouse Gases Tool (FLIGHT) system (<http://ghgdata.epa.gov/ghgp/main.do>) and data for individual facilities can be downloaded from that system. Data from reporting year 2012 was the latest data available at the time this work was completed. FLIGHT data from reporting year 2012 shows 58,663 wells that report unloading emissions. Of these 58,663 wells from FLIGHT, 32,225 are reported as having plunger lifts and 26,438 are reported as not having plunger lifts. This agrees reasonably well with the EPA NEI estimates for total well count (58,663 from FLIGHT as compared to 60,810 from the GHG NEI), however, the fractions of plunger and non-plunger wells differ for the two data sets. In this work, the fractions of plunger and non-plunger wells from the 2012 GHGRP will be used to

estimate national emissions, based on the assumption that these data are based on more recent assessments of the prevalence of plunger lifts. A sensitivity analysis is presented at the end of this section to assess the effect of this assumption on national emission estimates.

While this work uses total well counts from the 2012 GHGRP FLIGHT data in estimating national emissions, event counts will be based on data from a survey of companies participating in this work. One reason for using survey data for event counts is the lack of complete data on event frequency for non-plunger wells in the 2012 GHGRP FLIGHT data. While it may be feasible to estimate these data from partial reporting in the 2012 GHGRP, this would require making assumptions regarding the representativeness of partial reporting. A second reason for using the survey data collected in this work, rather than GHGRP FLIGHT data for event counts, was apparent quality assurance issues in event counts in the GHGRP FLIGHT data. Specifically, the study team's interpretation of the event reports in the GHGRP suggested very high frequencies of events for some non-plunger wells. These event frequencies appeared to the study team to be physically unreasonable (thousands of events per year for wells that are almost exclusively manually unloaded) and inconsistent with the survey data from companies.

Since event counts for plunger and non-plunger wells are either not reported or inconsistently reported in the FLIGHT data, the companies participating in this study volunteered data on unloading emissions that they had released to the EPA GHGRP for reporting year 2013. Eight of the ten participants were able to provide data. The participant company data, underwent quality assurance review and was then used to determine average event frequencies for the categories of non-plunger and plunger lift wells. The results for non-plunger lift wells are shown in Tables B-12 (national totals) and B-13 (regional distributions). The data in Table S5-3 indicate that event count distributions in the four regions considered in this work are all similar to national averages, but since detailed regional data are available, they will be used in estimating emission event counts in this work.

As an example of how event counts were estimated in Table B-12, consider the national total for non-plunger wells with less than 10 events per year. In the survey 6,378 of the 7,481 wells without plunger lift (85%, see Table B-12) had less than 10 events per year. It therefore is assumed that 85% of the 26,438 wells without plunger lift nationally will have less than 10 events per year, with an average of 2.93 events per well (see Table B-13, B-14). This results in an estimate of 66,000 events for these low event frequency wells in Table B-12. A total event count for all wells without plunger lifts reported through the GHGRP is estimated as 170,000 events, based on national average data, as shown in Table S5-2. If the averaging is done on a regional basis, as shown in Table S5-4, the national event count is estimated as 177,000 events. The regional estimate of event counts was used to produce a national methane emission estimate of 4.4 bcf/yr, as shown in Table B-15.

Table B-12. Activity data for wells without plunger lifts, based on surveys of participating companies using nationally averaged data

Well Type Strata	Total Number of events	% of events provided by participants	Number of Venting Wells	% of wells provided by participants	National Event Count if Participant Event Counts are scaled up by number of wells
Non Plunger Wells					
(events<10)	18,691	39	6,378	85	66,000
Non Plunger Wells					
(10≤events<50)	20,593	43	1,016	14	73,000
Non Plunger Wells					
(50≤events<200)	5,969	12	79	1	21,000
Non Plunger Wells					
(events≥201)	2,705	6	8	0.1	9,600
Total	47,958	100	7,481	100	170,000

Table B-13. Regional distributions of event counts, based on surveys of participating companies.

Company Data Well type (Annual Frequency of events)	AP			GC			MC			RM			Total		
	# of wells	#of events	Events/ Well	# of wells	#of events	Events/ Well	# of wells	#of events	Events/ Well	# of wells	#of events	Events/ Well	# of wells	#of events	Events/ Well
Non Plunger lift wells															
0-10	744	2,327	3.13	1,520	4,455	2.93	2,315	6,831	2.95	1,799	5,078	2.82	6,378	18,691	2.93
11-50	179	3,826	21.4	180	3,491	19.4	461	9,757	21.1	196	3,519	18.0	1,016	20,593	20.3
51-200	19	1,270	66.8	11	915	83.2	41	3,177	77.5	8	607	75.9	79	5,969	75.6
201+	0	0	-	1	355	355	7	2,350	336	0	0	-	8	2,705	338
Plunger lift wells															
0-99	42	302	7.19	423	2,237	5.29	857	4,419	5.16	3,845	60,282	15.7	5,167	67,240	13.01
100+	1	259	259	3	366	122	191	324,341	1,698	3,410	3,508,080	1,029	3,605	3,833,046	1,063

Table B-14. Regional distributions of non-plunger well event counts, based on surveys of participating companies.

Company Data Well type (Annual Frequency of events)	AP				GC				MC				RM				National event count
	# of wells	% of wells	Events/ Well	Region Event count	# of wells	% of wells	Events/ Well	Region Event count	# of wells	% of wells	Events/ Well	Region Event count	# of wells	% of wells	Events/ Well	Region Event count	
0-10	7812	79%	3.13	19,300	3855	88.8%	2.93	10,000	8219	82.0%	2.95	19,900	6552	89.8%	2.82	16,600	65,800
11-50		19%	21.4	31,700		10.5%	19.4	7,900		16.3%	21.1	28,400		9.8%	18.0	11,500	79,500
51-200		2%	66.8	10,500		0.6%	83.2	2,100		1.5%	77.5	9,200		0.4%	75.9	1,990	23,800
201+		0%	-	0		0.06%	355	800		0.2%	336	6,800		-	-	-	7,600
Total		100%		62,000		100%		21,000		100%		64,000		100%		30,000	177,000

Table B-15. National emission estimate for plunger lift wells

Well type (Annual Frequency of events)	Event count (events/yr)	Measured EF, scf methane per event (95% confidence range)	Emissions (billion scf methane/yr)
Non Plunger Wells (events<10)	65,800	21,500 (9,600-37,800)	1.4 (0.6-5.2)
Non Plunger Wells (10≤events<50)	79,500	24,100(8,700-50,400)	1.9 (0.7-4.0)
Non Plunger Wells (50≤events<200)	23,800	35,000 (18,700-53,000)	0.8 (0.4-1.3)
Non Plunger Wells (events≥200)	7,600	Not measured, assume 35,000 (see main text) (18,700-53,000)	0.3 (0.1-0.4)
Total			4.4 (2.8-8.5*)

*Range assumes that emission factors for each frequency range are independent.

Table B-16. Regional distributions of plunger lift well counts, and high and low frequency wells used in making national emission estimates

Well type (Annual Frequency of events)	AP		GC		MC		RM Basin 580		RM excluding Basin 580	
	well count	f _{high or low} freq. events	well count	f _{high or low} freq. events	well count	f _{high or low} freq. events	well count	f _{high or low} events	well count	f _{high or low} freq. events
		0-99		10,869		0.98		1,048		0.993
100+	0.02	0.007	0.18		0.68	0.09				

National event counts for plunger wells were estimated with a slightly modified procedure. The modifications are necessary because the distribution of high event frequency wells (≥ 100 events/yr) is not uniform within the Rocky Mountain region. As shown in Table B-16 and Table B-17, Basin 580 in the Rocky Mountain region has a much higher fraction of high frequency plunger lift unloadings than the rest of the region. Therefore, the total number of events associated with high and low frequency wells in the Rocky Mountain region was estimated using two sub-regions.

As an example, the total number of plunger lift wells in Basin 580 of the Rocky Mountain region (from the GHGRP) was 5,041 (Table B-16). Data from the company participant survey (Table B-16) indicated that 68% of the plunger lift wells in Basin 580 had 100 or more events per year, leading to an estimate of 3,404 high frequency plunger lift wells in Basin 580 (Table B-17). Similar calculations were done to estimate the number of low and high frequency wells in each region and sub-region. The event estimates are shown in Table B-17, and suggest a total of 206,500 events/yr for low frequency wells and 6.56 million events/yr for high frequency wells. National emission estimates, based on 5 region averaging (AP, GC, MC, two RM sub-regions), are shown in Table B-18.

Table B-17. Activity data for wells with plunger lifts, based on surveys of participating companies

Well Type	National Total Number of wells	$N_{\text{low or high freq. events, } i}$ (Average number of events per well)	Scaled total Number of events
AP Plunger Wells (events<100)	10,616	7.19	76,300
GC Plunger Wells (events<100)	1,041	5.29	5,500
MC Plunger Wells (events<100)	4,167	5.16	21,500
RM Plunger Wells (events<100)	10,930		103,200
(Basin 580 only)	(1,637)	(31.2)	(51,100)
[RM without basin 580]	[9,293]	[5.61]	[52,100]
Total Plunger Wells (events<100) (5 regions)	26,754		206,500
AP Plunger Wells (events 100+)	253	259	65,500
GC Plunger Wells (events 100+)	7	122	900
MC Plunger Wells (events 100+)	929	1,698	1,577,400
RM Plunger Wells (events 100+)	4,282		4,919,400
(Basin 580 only)	(3,404)	(976)	(3,322,300)
[RM without basin 580]	[878]	[1,819]	[1,597,100]
Total Plunger Wells (events 100+) (5 regions)	5,471		6,563,200
National Total (5 regions)	32,225		6,769,700

Table B-18. National emission estimate for plunger lift wells

Well type (Annual Frequency of events)	Event count, events/yr	Measured EF, scf methane per event (95% confidence range)	Emissions, billion scf methane/yr (95% confidence range)
Plunger Wells (events<100)	26,500	9,450 (6,900-12,400)	2.0 (1.4-2.6)
Plunger Wells (events≥100)	6,563,000	1,200 (500-2,100)	7.9 (3.3-13.8)
Total			9.9 (5.2-15.8*)

*Range assumes that emission factors for each frequency range are independent.

Taken together, methane emissions from both plunger lift wells (Table S5-8) and non-plunger lift wells (Table S5-5) are 14.3 billion scf/yr (270 Gg). The 95% confidence range, assuming that the estimates for plunger lift wells and non-plunger lift wells are independent, is 10-21 billion scf/yr (190-400 Gg/yr).

A sensitivity analysis on national emission estimates was performed using alternative distributions of plunger and non-plunger wells in the national well count. The national emission estimates reported in Tables B-15 and B-18 are based on total counts of plunger and non-plunger lift wells from the 2012 GHGRP. If the data from the 2012 GHG NEI are used instead, the total emission estimates would scale based on the number of wells in each category. So, if the total plunger well count is reduced from 32,225 to 23,503, total estimated emissions from plunger lift wells would decrease to 7.2 bcf ($9.9 \text{ bcf} * 23,503/32,225$). This decrease in plunger lift well emissions would be partially offset by increases in emissions from non-plunger wells. The count of plunger lift wells would increase from 26,438 to 37,307 and total estimated emissions from the non-plunger wells would increase to 6.2 bcf ($4.4 \text{ bcf} * 37,307/26,438$). Overall, emissions from both plunger lift and non-plunger lift wells would decrease slightly from 14 bcf/yr to 13 bcf/yr.

B.6 Comparison between measurements reported in this work and measurements reported by Allen et al. [1]

Allen *et al.* [1] reported measurements for 9 unloading events. All were manual unloadings of relatively recent horizontal wells, without plunger lifts, in newly developed shale formations. A comparison of the data reported by Allen *et al.* and the measurements reported in this work is provided in Table B-19.

Table B-19. Comparison of measurements from Allen, *et al.* [1] and similar measurements made in this work.

	Allen, <i>et al.</i> [1]	Measurements made in this work for horizontal wells without plunger lifts (see horizontal wells in Table B-8)
Average emissions per event (scf/event)	57,000	52,000
Range of emissions per event (scf/event)	950-191,000	1,400-135,000
Average frequency of unloadings (events/yr)	5.9	40.
Range of frequencies (events/yr)	1-12	1-101
Duration of event (hr)	1.0	2.2
Range of durations (hr)	0.25-2.77	0.35-4.5

The measurements from Allen, *et al.* [1] were not combined with the measurements in this work for two reasons:

1. Some of the wells on which Allen, *et al.* [1] made measurements were shut-in for a period of a week or more while the study team made arrangements to get to the

- site. This shut-in period was not part of routine unloading practices and may introduce a bias into the measurements.
2. If the measurements of Allen, *et al.* [1] were added to the results from Table B-8, more than half of the wells with manual unloadings would be horizontal wells. This may not accurately represent national populations.

B.7 References

(1) Allen DT, Torres VM, Thomas J, Sullivan D, Harrison M, Hendler, A., Herndon, SC, Kolb, CE, Fraser, MP, Hill, AD, *et al.*(2013) Measurements of Methane Emissions at Natural Gas Production Sites in the United States, *Proc. Natl. Acad. Sci. U.S.A.*, 110: 17768-17773.

C Analysis of Methane Emissions Attributable to Natural Gas Production

C.1 Data used to allocate emissions on an energy and price basis

Table C-1. Energy and Price Data

	Energy (HHV)*	Energy (LHV)	Price[4]
CH₄	1,010 BTU/scf	913 BTU/scf**	4 \$/Thousand scf
C₂H₆	1,769 BTU/scf	1,641 BTU/scf**	11 \$/Million BTU
C₃H₈	2,518 BTU/scf	2,385 BTU/scf**	11 \$/Million BTU
C₄H₁₀	3,262 BTU/scf	3,109 BTU/scf**	11 \$/Million BTU
C₅+	4,010 BTU/scf	3,882 BTU/scf**	11 \$/Million BTU
Hydrocarbon Liquids	5.8 Million BTU/bbl	5.4 Million BTU/bbl***	100 \$/bbl
Natural Gas	1,027 BTU/scf	983 BTU/scf***	4 \$/Thousand scf

*Table 3-7 and Table 3-8. Higher Heating Value (HHV) is used in this work [1].

** Adapted from Fuel Flue Gases, American Gas Association [2].

*** Adapted from GREET, Argonne [3].

C.2 Estimating rate of decrease in production over time and ultimate recovery (production)

The estimated ultimate recovery (EUR) for each of the sites is based on a decline curve analysis (DCA)[5] and historical data for shale plays. To determine production of the well (q_t) at time t , based on DCA:

$$q_t = \frac{q_i}{(1 + D_i \times b \times t)^{\frac{1}{b}}} \quad (\text{C2.1})$$

where:

t = age of well

q_t = Production of well at time t .

q_i = Initial production rate

D_i = initial decline rate

b = decline exponent

O'Sullivan and Paltsev [6] presented data on the rate of decrease of production from horizontal wells from various plays. Data were fitted to equation (C2.1) to determine the value of the two parameters; $D_i = 0.1264$, and $b = 1.2187$ (standard error = 0.0205, $r = 0.9984$).

For each production site where measurements were made by Allen, *et al* [8], the age of the wells (t) is provided, as well as the gas and oil production at the time of the measurements (q_t).

To find the initial production (q_i), from equation (C2.1):

$$q_i = q_t (1 + 0.1264 \times 1.2187 \times t)^{\frac{1}{1.2187}} \quad (\text{C2.2})$$

To find the EUR, from equation (C2.1) and (C2.2):

$$EUR = \int_0^{t_f} q_t dt \quad (\text{C2.3})$$

$$EUR = \int_0^{t_f} \frac{q_i}{(1 + 0.1264 \times 1.2187 \times t)^{\frac{1}{1.2187}}} dt \quad (\text{C2.4})$$

Where t_f is the well lifetime,

$$EUR = 36.1746 q_i \left[(0.1540 t_f + 1)^{0.1794} - 1 \right] \quad (\text{C2.5})$$

A well lifetime (t_f) of 30 years is assumed for all production sites. A sensitivity test was done, assuming a well lifetime of 15 years. Table C-2 shows gas and oil production at each well at the time of the measurement and the estimated ultimate production.

Table C-2. Gas and oil production at the time of the measurement (q_t), estimated initial production rate (q_i), and estimated ultimate recovery (EUR) at each site, assuming well lifetimes of 30 and 15 years. *The two letter code for each site indicates the region in which sampling was done (AP=Appalachia; GC=Gulf Coast; MC=Midcontinent; RM=Rocky Mountains)*

Site	Age of well (years)	Gas q_t (MMscf/day)	Oil q_t (bbl/day)	Gas q_i (MMscf/yr)	Oil q_i (bbl/yr)	Gas EUR (MMscf) 30-yr	Oil EUR (bbl) 30-yr	Gas EUR (MMscf) 15-yr	Oil EUR (bbl) 15-yr
AP-1	1.0	1.10	0	948	0	3,035	0	2,362	0
AP-2	0.5	5.42	0	3,385	0	10,839	0	8,436	0
AP-3	0.8	12.0	0	9,225	0	29,541	0	22,992	0
AP-4	2.5	8.95	0	13,471	0	43,137	0	33,574	0
AP-5	0.1	5.20	0	2,181	0	6,985	0	5,437	0
AP-6	2.7	5.21	0	8,263	0	26,459	0	20,593	0
AP-7	2.7	5.73	0	9,087	0	29,099	0	22,648	0
AP-8	0.8	1.44	3.36	1,107	2,583	3,545	8,271	2,759	6,461
AP-9	3.0	4.29	0	7,316	0	23,428	0	18,234	0
AP-10	3.0	5.80	0	9,891	0	31,674	0	24,652	0
AP-11	1.1	2.00	0	1,815	0	5,811	0	4,522	0
AP-12	2.0	1.96	0	2,546	0	8,152	0	6,345	0
AP-13	2.7	6.05	0	9,595	0	30,724	0	23,913	0
AP-14	2.0	4.75	0	6,170	0	19,756	0	15,376	0
AP-15	3.5	1.56	0	2,965	0	9,495	0	7,390	0
AP-16	2.2	1.48	0	2,046	0	6,551	0	5,098	0
AP-17	0.5	4.53	0	2,829	0	9,059	0	7,051	0
AP-18	0.4	27.4	0	15,751	0	50,438	0	39,256	0
AP-19	8.8	1.73	0	6,540	0	20,941	0	16,299	0
AP-20	2.0	3.79	0	4,923	0	15,764	0	12,269	0
AP-21	0.2	5.00	0	2,363	0	7,565	0	5,888	0
AP-22	1.5	22.5	0	24,383	0	78,078	0	60,768	0
AP-23	9.4	2.13	0	8,474	0	27,135	0	21,119	0
AP-24	2.7	31.8	0	50,432	0	161,494	0	125,690	0
AP-25	1.0	4.06	0	3,498	0	11,202	0	8,719	0
AP-26	0.7	12.0	0	8,657	0	27,721	0	21,575	0
AP-27	1.0	5.25	0	4,524	0	14,486	0	11,274	0
AP-28	0.6	2.50	0	1,683	0	5,390	0	4,195	0
AP-29	2.8	7.60	0	12,357	0	39,571	0	30,798	0
AP-31	0.7	12.0	0	8,657	0	27,721	0	21,575	0
AP-32	0.6	14.4	0	9,696	0	31,049	0	24,165	0
AP-33	1.0	26.3	0	22,636	0	72,484	0	56,414	0

Table C-2 (continued).

Site	Age of well (years)	Gas q _t (MMscf/day)	Oil q _t (bbl/day)	Gas q _i (MMscf/yr)	Oil q _i (bbl/yr)	Gas EUR (MMscf) 30-yr	Oil EUR (bbl) 30-yr	Gas EUR (MMscf) 15-yr	Oil EUR (bbl) 15-yr
AP-34	1.5	14.4	0	15,593	0	49,933	0	38,862	0
AP-35	1.0	2.63	0	2,266	0	7,257	0	5,648	0
AP-36	0.5	1.60	0	999	0	3,200	0	2,490	0
AP-37	0.8	7.20	0	5,535	0	17,725	0	13,795	0
AP-38	0.5	1.39	0	868	0	2,780	0	2,164	0
AP-39	1.8	19.4	0	23,571	0	75,479	0	58,745	0
AP-40	0.6	5.15	0	3,468	0	11,104	0	8,642	0
AP-41	0.5	11.2	0	6,995	0	22,398	0	17,433	0
AP-42	2.7	11.3	0	17,889	0	57,285	0	44,585	0
AP-43	0.3	10.7	0	5,625	0	18,012	0	14,019	0
AP-45	2.7	2.11	0	3,346	0	10,715	0	8,340	0
AP-46	1.0	8.00	0	6,893	0	22,074	0	17,180	0
AP-47	1.3	10.3	0	10,220	0	32,726	0	25,471	0
GC-1	1.0	0.23	273	198	234,905	635	752,215	494	587,544
GC-2	1.5	3.76	1,175	4,080	1,275,309	13,065	4,083,808	10,169	3,189,806
GC-3	0.3	0.89	0	467	0	1,494	0	1,163	0
GC-4	1.5	9.30	1,045	10,092	1,134,428	32,315	3,632,677	25,151	2,837,434
GC-5	1.0	3.27	1,163	2,818	1,002,322	9,023	3,209,646	7,022	2,507,010
GC-6	1.5	4.75	454	5,154	493,122	16,505	1,579,080	12,846	1,233,398
GC-7	1.6	1.45	0	1,636	0	5,240	0	4,078	0
GC-8	3.2	0.79	0	1,409	0	4,513	0	3,513	0
GC-9	1.5	13.4	950	14,551	1,030,561	46,597	3,300,071	36,266	2,577,640
GC-10	0.4	44.0	0	25,311	0	81,050	0	63,081	0
GC-11	2.5	5.75	0	8,655	0	27,714	0	21,570	0
GC-12	2.7	11.8	0	18,714	0	59,925	0	46,640	0
GC-13	2.1	1.78	0	2,386	0	7,642	0	5,948	0
GC-14	1.5	4.71	919	5,111	997,410	16,366	3,193,916	12,738	2,494,724
GC-15	1.4	19.3	0	20,045	0	64,188	0	49,958	0
GC-16	0.6	4.22	2,946	2,841	1,983,425	9,099	6,351,344	7,082	4,960,946
GC-17	0.3	33.2	0	17,378	0	55,648	0	43,311	0
GC-18	0.3	32.8	0	17,210	0	55,111	0	42,893	0
GC-19	1.4	15.3	0	15,911	0	50,950	0	39,655	0
GC-20	2.7	0.69	0	1,094	0	3,504	0	2,727	0
GC-21	1.8	1.20	0	1,457	0	4,666	0	3,632	0
GC-22	4.4	0.50	0	1,121	0	3,589	0	2,793	0
GC-23	0.4	12.8	0	7,335	0	23,489	0	18,281	0

Table C-2 (continued).

Site	Age of well (years)	Gas q_t (MMscf/day)	Oil q_t (bbl/day)	Gas q_i (MMscf/yr)	Oil q_i (bbl/yr)	Gas EUR (MMscf) 30-yr	Oil EUR (bbl) 30-yr	Gas EUR (MMscf) 15-yr	Oil EUR (bbl) 15-yr
GC-24	1.5	3.00	538	3,255	583,968	10,424	1,869,989	8,113	1,460,623
GC-25	8.7	0.03	0	112	0	360	0	280	0
GC-26	1.5	13.9	508	15,072	551,209	48,265	1,765,086	37,564	1,378,684
GC-27	10	0.09	0	382	0	1,222	0	951	0
GC-28	1.6	11.8	0	13,283	0	42,535	0	33,105	0
GC-29	2.0	0.66	0	857	0	2,745	0	2,137	0
GC-30	1.1	5.99	0	5,435	0	17,403	0	13,545	0
GC-31	0.7	6.51	1,833	4,696	1,322,018	15,039	4,233,380	11,704	3,306,634
GC-32	5.9	2.37	0	6,604	0	21,146	0	16,458	0
GC-33	4.2	0.82	0	1,777	0	5,690	0	4,428	0
GC-34	0.4	37.3	0	21,413	0	68,570	0	53,368	0
GC-35	5.9	1.97	0	5,489	0	17,577	0	13,680	0
GC-36	0.2	0.83	0	392	0	1,256	0	977	0
GC-37	1.5	6.01	1,163	6,522	1,262,266	20,883	4,042,042	16,254	3,157,183
GC-38	1.6	1.79	0	2,020	0	6,469	0	5,035	0
GC-39	5.6	0.67	0	1,795	0	5,749	0	4,474	0
GC-40	2.6	0.18	0	278	0	891	0	693	0
GC-41	6.4	0.04	0	119	0	379	0	295	0
GC-42	0.3	1.43	0	750	0	2,401	0	1,868	0
GC-43	8.7	0.10	0	375	0	1,200	0	934	0
GC-44	8.5	0.03	0	110	0	354	0	275	0
GC-45	9.9	0.02	0	83	0	265	0	206	0
GC-46	8.4	0.36	1.16	1,313	4,230	4,204	13,546	3,272	10,581
GC-47	9.0	0.06	0	231	0	739	0	575	0
GC-48	3.3	0.23	0	419	0	1,343	0	1,045	0
GC-49	9.9	0.05	0	207	0	663	0	516	0
GC-50	3.0	3.36	0	5,730	0	18,349	0	14,281	0
GC-51	9.5	0.07	0	281	0	899	0	700	0
GC-52	1.0	14.6	0	12,554	0	40,202	0	31,289	0
GC-53	0.5	1.73	0	1,080	0	3,460	0	2,693	0
GC-54	0.5	3.16	987	1,973	616,608	6,320	1,974,509	4,918	1,542,261
GC-55	0.9	47.7	0	38,891	0	124,538	0	96,927	0
GC-56	3.5	0.94	0	1,787	0	5,721	0	4,453	0
GC-57	1.2	15.5	0	14,762	0	47,272	0	36,792	0
GC-58	7.1	0.92	0	2,949	0	9,444	0	7,350	0
MC-1	0.9	0.88	47.1	718	38,426	2,298	123,049	1,789	96,112

Table C-2 (continued).

Site	Age of well (years)	Gas q_t (MMscf/day)	Oil q_t (bbl/day)	Gas q_i (MMscf/yr)	Oil q_i (bbl/yr)	Gas EUR (MMscf) 30-yr	Oil EUR (bbl) 30-yr	Gas EUR (MMscf) 15-yr	Oil EUR (bbl) 15-yr
MC-4	2.0	3.15	0	4,091	0	13,102	0	10,197	0
MC-5	0.0	0.03	0	11	0	35	0	27	0
MC-6	1.1	1.47	30.1	1,334	27,273	4,271	87,333	3,324	68,215
MC-7	3.0	0.10	0	171	0	546	0	425	0
MC-8	5.0	0.61	0	1,502	0	4,810	0	3,744	0
MC-9	3.0	0.21	0	358	0	1,147	0	893	0
MC-10	2.0	2.08	0	2,702	0	8,651	0	6,733	0
MC-11	4.0	0.98	0	2,050	0	6,564	0	5,108	0
MC-12	0.3	0.61	79.6	320	41,702	1,024	133,538	797	104,305
MC-13	4.0	0.43	0	899	0	2,880	0	2,241	0
MC-14	0.7	0.28	0	202	0	647	0	503	0
MC-16	1.1	0.91	32.4	826	29,432	2,644	94,248	2,058	73,616
MC-17	2.0	4.06	0	5,273	0	16,887	0	13,143	0
MC-18	4.0	0.66	0	1,380	0	4,420	0	3,440	0
MC-19	4.0	1.39	0	2,907	0	9,310	0	7,246	0
MC-20	0.8	2.07	43.3	1,591	33,311	5,096	106,667	3,966	83,316
MC-21	0.6	0.71	155	478	104,583	1,531	334,897	1,191	261,584
MC-22	5.0	2.90	0	7,142	0	22,869	0	17,799	0
MC-23	2.0	6.88	0	8,936	0	28,616	0	22,272	0
MC-24	5.0	0.25	0	616	0	1,971	0	1,534	0
MC-25	2.0	1.70	0	2,208	0	7,071	0	5,503	0
MC-26	5.0	0.22	0	542	0	1,735	0	1,350	0
RM-2	5.9	0.14	7.54	390	21,009	1,249	67,275	972	52,548
RM-3	4.0	0.05	1.73	105	3,618	335	11,587	261	9,050
RM-4	1.0	0.42	70.0	362	60,333	1,159	193,200	902	150,906
RM-5	3.8	0.17	8.56	343	17,254	1,097	55,251	854	43,156
RM-6	2.5	0.10	2.43	151	3,658	482	11,712	375	9,148
RM-7	7.0	0.05	1.75	159	5,550	508	17,771	395	13,881
RM-8	0.0	0.50	96.7	183	35,299	584	113,035	455	88,290
RM-9	9.0	0.06	2.59	231	9,962	739	31,901	575	24,918
RM-10	3.3	0.18	11.4	328	20,820	1,051	66,669	818	52,074
RM-11	2.0	0.20	2.09	260	2,715	832	8,693	647	6,790
RM-12	1.0	0.07	8.76	60	7,548	193	24,171	150	18,879
RM-13	7.0	0.11	22.9	349	72,589	1,117	232,445	869	181,560
RM-14	5.0	0.23	7.53	566	18,544	1,814	59,380	1,412	46,381
RM-15	8.0	0.29	40.5	1,019	142,251	3,262	455,519	2,539	355,800

Table C-2 (continued).

Site	Age of well (years)	Gas q_t (MMscf/day)	Oil q_t (bbl/day)	Gas q_i (MMscf/yr)	Oil q_i (bbl/yr)	Gas EUR (MMscf) 30-yr	Oil EUR (bbl) 30-yr	Gas EUR (MMscf) 15-yr	Oil EUR (bbl) 15-yr
RM-16	2.5	0.03	1.63	45	2,453	145	7,856	113	6,136
RM-17	1.0	2.58	81.0	2,223	69,768	7,119	223,413	5,541	174,505
RM-18	1.0	0.32	21.6	276	18,595	883	59,544	687	46,509
RM-19	5.0	0.23	7.53	566	18,544	1,814	59,380	1,412	46,381

C.3 Total Routine Emissions measured by site

Table C-3. Measured methane emissions, assuming a constant emission rate, and a 30-yr well lifetime. Last column represents the ratio of total emissions at each site, over the 30 year period, divided by the estimated gas ultimate recovery; assuming 30-yr well lifetime.

Methane Emissions over a 30 year period (Mg)						
Site	Wells	Pneumatic Controllers	Pneumatic Pumps	Equipment Leaks	Total	Total Emissions (Mg) / Gas EUR (Mg) (30- yr)
AP-1	1	38.15	0	0	38.15	0.0006
AP-2	3	41.17	0	0	41.17	0.0002
AP-3	4	259.4	0	3.027	262.4	0.0005
AP-4	4	43.60	0	46.62	90.22	0.0001
AP-5	6	5.449	0	26.04	31.49	0.0002
AP-6	5	8.477	58.13	0	66.60	0.0001
AP-7	3	106.0	58.13	0	164.1	0.0003
AP-8	1	168.5	0	0	168.5	0.0019
AP-9	4	148.0	0	522.8	670.9	0.0014
AP-10	5	304.6	0	370.0	674.5	0.0011
AP-11	1	11.35	0	0	11.35	9.9E-05
AP-12	1	33.30	0	86.59	119.9	0.0007
AP-13	3	240.4	58.13	0	298.5	0.0005
AP-14	6	70.54	0	0	70.54	0.0002
AP-15	2	0	0	0	0	0
AP-16	1	65.39	0	0	65.39	0.0005
AP-17	2	95.06	0	258.2	353.3	0.0020
AP-18	6	5.752	0	81.74	87.49	8.8E-05
AP-19	3	86.59	0	29.37	116.0	0.0003
AP-20	1	0	0	0	0	0
AP-21	1	106.0	0	47.23	153.2	0.0010
AP-22	3	136.2	174.4	0	310.6	0.0002
AP-23	6	125.9	58.13	136.5	320.6	0.0006
AP-24	2	48.14	58.13	0	106.3	3.3E-05
AP-25	5	104.4	0	1,653	1,758	0.0080
AP-26	3	174.4	0	20.89	195.3	0.0004
AP-27	7	335.4	0	512.5	848.0	0.0030
AP-28	8	43.60	0	0	43.60	0.0004
AP-29	3	73.26	0	90.52	163.8	0.0002
AP-31	5	231.6	0	23.31	254.9	0.0005
AP-32	5	51.47	0	47.23	98.70	0.0002
AP-33	7	1,811	348.8	4.238	2,164	0.0015

Table C-3 (continued).

Methane Emissions over a 30 year period (Mg)						
Site	Wells	Pneumatic Controllers	Pneumatic Pumps	Equipment Leaks	Total	Total Emissions (Mg) / Gas EUR (Mg) (30- yr)
AP-34	4	1,893	290.6	0.605	2,184	0.0022
AP-35	3	380.4	0	0	380.4	0.0027
AP-36	1	9.082	0	0	9.082	0.0001
AP-37	5	9.082	0	172.3	181.3	0.0005
AP-38	2	116.0	0	378.7	494.7	0.0092
AP-39	3	163.5	174.4	68.42	406.3	0.0003
AP-40	1	3.330	58.13	3.330	64.79	0.0003
AP-41	8	0	0	29.97	29.97	6.8E-05
AP-42	3	877.8	174.4	0	1,052	0.0009
AP-43	6	41.48	0	852.2	893.7	0.0025
AP-45	2	17.26	0	25.13	42.38	0.0002
AP-46	3	272.5	0	0	272.5	0.0006
AP-47	6	40.27	0	2.422	42.69	6.5E-05
GC-1	3	334.2	0	48.74	383.0	0.0231
GC-2	6	635.8	0	4.541	640.3	0.0018
GC-3	1	337.3	0	0	337.3	0.0123
GC-4	7	662.6	0	48.14	710.8	0.0008
GC-5	6	1,695	0	372.4	2,068	0.0084
GC-6	6	847.7	0	81.14	928.8	0.0023
GC-7	1	64.48	83.86	0	148.3	0.0015
GC-8	1	17.26	58.13	0	75.38	0.0009
GC-9	6	75.08	0	0	75.08	6.5E-05
GC-10	4	847.7	0	0	847.7	0.0006
GC-11	2	135.9	0	22.10	158.0	0.0003
GC-12	0	729.0	0	26.64	755.7	0.0007
GC-13	1	248.6	0	3.633	252.2	0.0018
GC-14	3	847.7	0	0	847.7	0.0017
GC-15	6	694.6	0	258.5	953.2	0.0008
GC-16	0	1,673	0	0	1,673	0.0075
GC-17	3	1,082	0	0	1,082	0.0011
GC-18	3	265.8	0	32.70	298.5	0.0003
GC-19	6	1,115	0	17.26	1,132	0.0012
GC-20	1	52.98	0	19.68	72.66	0.0011
GC-21	1	13.32	0	0	13.32	0.0002
GC-22	1	30.88	299.7	16.65	347.2	0.0051

Table C-3 (continued).

Methane Emissions over a 30 year period (Mg)						
Site	Wells	Pneumatic Controllers	Pneumatic Pumps	Equipment Leaks	Total	Total Emissions (Mg) / Gas EUR (Mg) (30- yr)
GC-23	2	45.41	0	0	45.41	0.0001
GC-24	5	635.8	0	27.25	663.0	0.0023
GC-25	1	91.73	115.0	16.65	223.4	0.0299
GC-26	5	779.9	62.67	69.93	912.5	0.0008
GC-27	1	79.92	93.25	0	173.2	0.0069
GC-28	5	185.7	0	27.55	213.2	0.0003
GC-29	1	100.5	0	92.64	193.2	0.0038
GC-30	3	94.46	286.7	98.09	479.2	0.0015
GC-31	0	218.0	0	0	218.0	0.0006
GC-32	1	106.0	188.3	0	294.3	0.0007
GC-33	1	52.98	86.28	0	139.3	0.0013
GC-34	3	845.6	0	0	845.6	0.0007
GC-35	1	317.9	1,301	0	1,618	0.0048
GC-36	1	607.9	0	0	607.9	0.0267
GC-37	16	50.86	0	57.52	108.4	0.0002
GC-38	1	61.15	58.13	0	119.3	0.0010
GC-39	1	106.0	1,442	42.38	1,591	0.0145
GC-40	1	156.5	0	24.22	180.7	0.0108
GC-41	1	640.3	75.08	35.72	751.1	0.1032
GC-42	1	534.6	0	0	534.6	0.0121
GC-43	1	71.15	58.13	0	129.3	0.0056
GC-44	1	4.238	0	0	4.238	0.0006
GC-45	1	106.9	0	0	106.9	0.0182
GC-46	1	138.1	0	0	138.1	0.0015
GC-47	1	77.81	212.8	0	290.6	0.0186
GC-48	1	57.22	0	90.52	147.7	0.0059
GC-49	1	65.09	7.871	215.9	288.8	0.0206
GC-50	2	34.51	58.13	51.47	144.1	0.0004
GC-51	1	142.3	179.5	0	321.8	0.0170
GC-52	3	358.1	182.3	0	540.4	0.0007
GC-53	1	455.9	0	0	455.9	0.0071
GC-54	0	50.05	0	10.90	60.95	0.0004
GC-55	6	535.9	58.13	0	594.0	0.0003
GC-56	1	2.422	116.3	13.93	132.6	0.0012
GC-57	5	134.4	0	93.25	227.7	0.0003

Table C-3 (continued).

Methane Emissions over a 30 year period (Mg)						
Site	Wells	Pneumatic Controllers	Pneumatic Pumps	Equipment Leaks	Total	Total Emissions (Mg) / Gas EUR (Mg) (30- yr)
GC-58	1	3.027	832.4	13.02	848.4	0.0047
MC-1	2	476.8	0.605	69.93	547.4	0.0097
MC-4	11	122.6	149.0	133.8	405.4	0.0015
MC-5	2	183.2	0	99.60	282.8	0.3548
MC-6	4	1,589	0	0	1,589	0.0165
MC-7	1	51.47	11.20	0	62.67	0.0060
MC-8	3	421.4	53.89	23.31	498.6	0.0053
MC-9	5	299.7	58.13	101.4	459.3	0.0207
MC-10	3	81.14	13.32	0	94.46	0.0006
MC-11	5	502.6	74.17	294.3	871.0	0.0069
MC-12	1	476.8	0	2.119	478.9	0.0188
MC-13	4	1,103	58.13	0	1,161	0.0207
MC-14	1	294.3	0	3.633	297.9	0.0197
MC-16	2	426.3	0	101.7	528.0	0.0086
MC-17	5	1,662	59.64	39.96	1,762	0.0054
MC-18	1	158.9	22.40	0	181.3	0.0021
MC-19	5	963.9	101.1	0	1,065	0.0059
MC-20	3	330.7	0	88.40	419.2	0.0036
MC-21	3	165.7	0	125.3	291.1	0.0081
MC-22	5	4,010	73.67	56.92	4,140	0.0093
MC-23	5	122.6	290.6	0	413.2	0.0008
MC-24	1	106.0	58.13	0	164.1	0.0043
MC-25	3	68.12	52.07	31.79	152.0	0.0011
MC-26	1	106.0	4.238	0	110.2	0.0033
RM-2	2	19.07	0	8.780	27.85	0.0008
RM-3	7	32.70	0	1.816	34.51	0.0039
RM-4	6	8.174	0	3.027	11.20	0.0004
RM-5	8	1,430	0	92.34	1,523	0.0526
RM-6	2	635.8	0	73.57	709.3	0.0560
RM-7	8	635.8	0	7.266	643.0	0.0533
RM-8	8	0	0	31.18	31.18	0.0021
RM-9	1	3.633	0	4.844	8.477	0.0004
RM-10	1	152.6	0	0.908	153.5	0.0055
RM-11	5	0	0	1.514	1.514	7.1E-05
RM-12	4	0	0	5.752	5.752	0.0011

Table C-3 (continued).

Methane Emissions over a 30 year period (Mg)						
Site	Wells	Pneumatic Controllers	Pneumatic Pumps	Equipment Leaks	Total	Total Emissions (Mg) / Gas EUR (Mg) (30- yr)
RM-13	7	0	0	0.605	0.605	2.1E-05
RM-14	1	635.8	0	21.19	657.0	0.0137
RM-15	9	0	0	0	0	0
RM-16	2	5.449	0	4.844	10.29	0.0027
RM-17	4	259.8	0	69.03	328.8	0.0017
RM-18	2	43.60	0	0	43.60	0.0019
RM-19	1	12.72	0	70.24	82.95	0.0018

C.4 Mass based allocation of emissions

To determine the fraction of non-methane hydrocarbon gases that are considered as part of *salable natural gas* (NG) and of *natural gas liquids* (NGL), a composition of 92.8% methane, 5.54% non-methane hydrocarbon (mass) is assumed for NG (the rest is N₂, CO₂, H₂S, and H₂O). This natural gas composition is based on frequently used life cycle datasets [7]. Consequently, if the gas produced at each site has a non-methane hydrocarbon content that would produce a *salable natural gas* of up to 5.54% non-methane hydrocarbon (mass) (Methane composition at that site would be 92.8% or higher (94.4% or higher for composition based on hydrocarbons only)), all non-methane hydrocarbon gases are considered as part of NG (for that site there would be no NGL production). On the other hand, if the gas produced at each site is wetter (methane composition at site would be less than 92.8%; 94.4% hydrocarbons only), non-methane hydrocarbon gases would represent more than 5.54% (5.6% hydrocarbon only), and they would be assigned to NG, starting with ethane and then adding subsequent heavier hydrocarbons, until the 5.54% threshold is reached, the remaining portion above 5.54% is considered as part of NGL.

Table C-4 shows the composition of salable natural gas and natural gas liquids (% mass) for each site. Table C-5 shows the fraction of production for each of the salable products (mass basis). Table C-6 shows the allocation of emissions from pneumatic controllers, pneumatic pumps, and equipment leaks over a 30 year period to each of the salable products, based. Table C-7 reports the ratio of allocated emissions to production for each of the three products: Natural Gas, Natural Gas Liquids, and Oil.

Table C-4. Composition of salable Natural Gas, and Natural Gas Liquids (% mass).

Site	Natural Gas					Natural Gas, EUR (Mg)	Natural Gas Liquids				Natural Gas Liquids, EUR (Mg)
	% C ₁	% C ₂	% C ₃	% C ₄	% C ₅₊		% C ₂	% C ₃	% C ₄	% C ₅₊	
AP-1	96.4	3.48	0.11	0	0	59,005	0	0	0	0	0
AP-2	95.5	4.19	0.27	0.04	0	211,289	0	0	0	0	0
AP-3	96.0	3.82	0.16	0.04	0	576,497	0	0	0	0	0
AP-4	97.2	2.76	0.08	0	0	836,980	0	0	0	0	0
AP-5	95.8	4.03	0.19	0	0	136,625	0	0	0	0	0
AP-6	95.7	4.04	0.24	0	0	516,860	0	0	0	0	0
AP-7	95.7	4.03	0.22	0	0	565,897	0	0	0	0	0
AP-8	94.4	5.60	0	0	0	55,351	49.3	28.60	13.78	8.34	32,392
AP-9	97.1	2.78	0.11	0.04	0	455,027	0	0	0	0	0
AP-10	97.0	2.89	0.11	0	0	614,983	0	0	0	0	0
AP-11	96.3	3.53	0.16	0.04	0	113,272	0	0	0	0	0
AP-12	94.4	5.23	0.03	0	0	160,036	0	91.4	8.65	0	656.0
AP-13	96.0	3.82	0.16	0	0	599,499	0	0	0	0	0
AP-14	94.9	4.68	0.32	0.07	0	387,376	0	0	0	0	0
AP-15	97.4	2.47	0.08	0.04	0	184,176	0	0	0	0	0
AP-16	94.7	4.87	0.35	0.04	0	128,401	0	0	0	0	0
AP-17	96.6	3.22	0.14	0.04	0	176,088	0	0	0	0	0
AP-18	95.1	4.61	0.27	0.04	0	988,031	0	0	0	0	0
AP-19	95.3	4.34	0.27	0.07	0	410,082	0	0	0	0	0
AP-20	97.2	2.68	0.11	0	0	305,353	0	0	0	0	0
AP-21	95.6	4.14	0.22	0.04	0	147,975	0	0	0	0	0
AP-22	96.0	3.82	0.16	0	0	1,523,474	0	0	0	0	0
AP-23	96.1	3.75	0.16	0	0	529,064	0	0	0	0	0
AP-24	95.7	4.03	0.22	0	0	3,140,582	0	0	0	0	0
AP-25	96.9	3.00	0.14	0	0	217,440	0	0	0	0	0
AP-26	96.0	3.82	0.16	0.04	0	540,977	0	0	0	0	0
AP-27	96.8	3.02	0.14	0	0	281,558	0	0	0	0	0
AP-28	96.0	3.82	0.16	0.04	0	105,196	0	0	0	0	0
AP-29	95.7	4.09	0.14	0.04	0	773,016	0	0	0	0	0
AP-31	96.0	3.82	0.16	0.04	0	540,977	0	0	0	0	0
AP-32	96.0	3.82	0.16	0.04	0	605,927	0	0	0	0	0
AP-33	95.6	4.16	0.24	0	0	1,417,505	0	0	0	0	0
AP-34	95.7	4.09	0.22	0	0	975,788	0	0	0	0	0
AP-35	95.6	4.04	0.24	0.04	0.10	141,472	0	0	0	0	0
AP-36	96.0	3.82	0.16	0.04	0	62,444	0	0	0	0	0

Table C-4 (continued).

Site	Natural Gas					Natural Gas, EUR (Mg)	Natural Gas Liquids				Natural Gas Liquids, EUR (Mg)
	% C ₁	% C ₂	% C ₃	% C ₄	% C ₅₊		% C ₂	% C ₃	% C ₄	% C ₅₊	
AP-37	95.1	4.61	0.27	0.04	0	347,208	0	0	0	0	0
AP-38	97.6	2.22	0.08	0.07	0	53,663	0	0	0	0	0
AP-39	95.4	4.34	0.24	0.04	0	1,477,859	0	0	0	0	0
AP-40	94.4	5.60	0	0	0	215,258	64.2	30.1	5.68	0	5,444
AP-41	96.0	3.82	0.16	0.04	0	437,108	0	0	0	0	0
AP-42	95.4	4.32	0.24	0.04	0	1,121,524	0	0	0	0	0
AP-43	95.1	4.61	0.27	0.04	0	352,842	0	0	0	0	0
AP-45	95.5	4.27	0.19	0	0	209,670	0	0	0	0	0
AP-46	96.1	3.65	0.16	0.07	0	430,507	0	0	0	0	0
AP-47	94.9	4.80	0.27	0	0	640,901	0	0	0	0	0
GC-1	94.4	5.60	0	0	0	9,665	38.8	30.5	17.7	13.0	6,725
GC-2	94.4	5.60	0	0	0	188,404	35.6	30.7	19.8	13.9	168,037
GC-3	99.8	0.16	0	0	0	27,566	0	0	0	0	0
GC-4	94.4	5.60	0	0	0	478,948	37.8	29.0	20.5	12.6	376,161
GC-5	94.4	5.60	0	0	0	132,917	36.2	28.9	19.8	15.1	108,584
GC-6	94.4	5.60	0	0	0	264,229	43.6	27.7	17.4	11.2	139,741
GC-7	99.8	0.16	0	0	0	96,687	0	0	0	0	0
GC-8	99.4	0.56	0.03	0	0	84,653	0	0	0	0	0
GC-9	94.4	5.60	0	0	0	746,434	43.5	27.8	17.4	11.2	394,305
GC-10	99.8	0.16	0	0	0	1,495,472	0	0	0	0	0
GC-11	99.7	0.30	0	0	0	506,093	0	0	0	0	0
GC-12	99.8	0.16	0	0	0	1,105,696	0	0	0	0	0
GC-13	99.7	0.28	0	0	0	139,183	0	0	0	0	0
GC-14	94.4	5.60	0	0	0	224,356	30.8	26.6	18.6	24.0	256,432
GC-15	99.7	0.31	0	0	0	1,176,216	0	0	0	0	0
GC-16	94.4	5.60	0	0	0	140,409	44.4	31.7	14.9	8.90	81,301
GC-17	99.8	0.16	0	0	0	1,028,161	0	0	0	0	0
GC-18	99.8	0.16	0	0	0	1,018,236	0	0	0	0	0
GC-19	99.6	0.33	0.03	0	0	938,788	0	0	0	0	0
GC-20	99.9	0.08	0	0	0	64,113	0	0	0	0	0
GC-21	99.8	0.16	0	0	0	86,101	0	0	0	0	0
GC-22	99.6	0.29	0.06	0	0	65,716	0	0	0	0	0
GC-23	99.8	0.16	0	0	0	433,391	0	0	0	0	0
GC-24	94.4	5.60	0	0	0	151,446	34.2	28.1	21.1	16.5	135,176
GC-25	94.4	5.60	0	0	0	6,787	16.1	32.5	20.5	30.9	415.3

Table C-4 (continued).

Site	Natural Gas					Natural Gas, EUR (Mg)	Natural Gas Liquids				Natural Gas Liquids, EUR (Mg)
	% C ₁	% C ₂	% C ₃	% C ₄	% C ₅₊		% C ₂	% C ₃	% C ₄	% C ₅₊	
GC-26	94.4	5.60	0	0	0	787,092	40.9	26.3	18.5	14.3	387,311
GC-27	94.4	5.60	0	0	0	23,124	0.6	37.1	23.6	38.7	1,044
GC-28	99.7	0.31	0	0	0	782,134	0	0	0	0	0
GC-29	99.7	0.24	0	0	0.06	50,244	0	0	0	0	0
GC-30	99.8	0.16	0	0	0	321,101	0	0	0	0	0
GC-31	94.4	5.60	0	0	0	248,763	42.4	28.6	16.9	12.2	109,222
GC-32	99.7	0.26	0.06	0	0	386,840	0	0	0	0	0
GC-33	99.5	0.38	0.09	0	0	103,660	0	0	0	0	0
GC-34	99.8	0.16	0	0	0	1,265,190	0	0	0	0	0
GC-35	99.8	0.22	0	0	0	323,297	0	0	0	0	0
GC-36	99.9	0.10	0	0	0	22,905	0	0	0	0	0
GC-37	94.4	5.60	0	0	0	299,193	33.1	27.1	19.2	20.6	282,666
GC-38	99.8	0.16	0	0	0	119,358	0	0	0	0	0
GC-39	97.8	1.32	0.31	0.19	0.36	106,442	0	0	0	0	0
GC-40	99.7	0.24	0	0	0.06	16,306	0	0	0	0	0
GC-41	97.8	1.32	0.31	0.19	0.36	7,027	0	0	0	0	0
GC-42	99.8	0.16	0	0	0	44,292	0	0	0	0	0
GC-43	97.8	1.32	0.31	0.19	0.36	22,216	0	0	0	0	0
GC-44	94.4	5.60	0	0	0	6,629	1.50	22.2	19.6	56.7	714.6
GC-45	94.4	5.60	0	0	0	4,850	29.9	24.5	16.1	29.5	788.4
GC-46	94.4	5.60	0	0	0	76,074	36.2	23.3	13.2	27.2	13,254
GC-47	94.4	5.60	0	0	0	13,820	15.6	29.3	21.0	34.0	1,197
GC-48	99.7	0.28	0	0	0	24,479	0	0	0	0	0
GC-49	94.4	5.60	0	0	0	12,400	15.6	29.3	21.0	34.0	1,074
GC-50	98.2	1.41	0.20	0.15	0.10	347,560	0	0	0	0	0
GC-51	94.4	5.60	0	0	0	16,828	12.3	26.9	21.1	39.7	1,394
GC-52	99.7	0.24	0	0	0.06	735,815	0	0	0	0	0
GC-53	99.8	0.16	0	0	0	63,836	0	0	0	0	0
GC-54	94.4	5.60	0	0	0	106,335	45.1	28.1	15.6	11.2	39,993
GC-55	99.9	0.10	0	0	0	2,273,091	0	0	0	0	0
GC-56	99.1	0.85	0.06	0	0	107,109	0	0	0	0	0
GC-57	99.6	0.33	0	0	0.06	868,706	0	0	0	0	0
GC-58	97.8	1.32	0.31	0.19	0.36	174,867	0	0	0	0	0
MC-1	94.4	5.60	0	0	0	33,134	29.0	37.6	19.5	13.8	22,571
MC-4	94.4	5.60	0	0	0	249,100	58.8	31.6	8.11	1.44	7,872

Table C-4 (continued).

Site	Natural Gas					Natural Gas, EUR (Mg)	Natural Gas Liquids				Natural Gas Liquids, EUR (Mg)
	% C ₁	% C ₂	% C ₃	% C ₄	% C ₅₊		% C ₂	% C ₃	% C ₄	% C ₅₊	
MC-5	94.4	5.60	0	0	0	529	36.2	32.9	15.8	15.0	281.6
MC-6	94.4	5.60	0	0	0	66,789	32.5	36.0	16.8	14.7	28,134
MC-7	98.7	1.22	0.06	0	0	10,182	0	0	0	0	0
MC-8	97.8	2.13	0.08	0	0	90,266	0	0	0	0	0
MC-9	97.3	2.66	0.06	0	0	21,572	0	0	0	0	0
MC-10	98.1	1.83	0.08	0	0	162,090	0	0	0	0	0
MC-11	97.7	2.13	0.08	0	0.06	123,083	0	0	0	0	0
MC-12	94.4	5.60	0	0	0	14,956	30.5	36.3	19.4	13.7	10,493
MC-13	97.3	2.61	0.06	0	0	54,485	0	0	0	0	0
MC-14	94.4	5.60	0	0	0	10,271	37.5	33.2	16.9	12.5	4,745
MC-16	94.4	5.60	0	0	0	42,652	36.8	34.6	16.0	12.7	18,094
MC-17	97.8	2.13	0.08	0	0	316,873	0	0	0	0	0
MC-18	96.9	3.10	0	0	0	83,731	0	0	0	0	0
MC-19	97.1	2.81	0.11	0	0	175,342	0	0	0	0	0
MC-20	94.4	5.60	0	0	0	80,002	36.6	35.9	15.7	11.7	35,579
MC-21	94.4	5.60	0	0	0	24,987	34.8	33.7	17.2	14.2	10,803
MC-22	97.2	2.69	0.11	0	0	432,029	0	0	0	0	0
MC-23	98.3	1.60	0.06	0	0.06	535,876	0	0	0	0	0
MC-24	97.8	2.12	0.06	0	0	36,863	0	0	0	0	0
MC-25	98.2	1.73	0.06	0	0	132,448	0	0	0	0	0
MC-26	97.8	2.11	0.06	0	0	32,519	0	0	0	0	0
RM-2	94.4	5.60	0	0	0	18,938	33.9	25.4	16.7	24.0	13,074
RM-3	94.4	5.60	0	0	0	5,104	34.6	29.7	21.0	14.7	3,410
RM-4	94.4	5.60	0	0	0	17,958	34.6	25.6	17.9	22.0	11,015
RM-5	94.4	5.60	0	0	0	16,636	33.9	25.4	16.7	24.0	11,484
RM-6	94.4	5.60	0	0	0	7,307	33.9	25.4	16.7	24.0	5,044
RM-7	94.4	5.60	0	0	0	8,457	40.6	23.6	13.5	22.3	3,182
RM-8	94.4	5.60	0	0	0	8,860	33.9	25.4	16.7	24.0	6,116
RM-9	94.4	5.60	0	0	0	11,204	33.9	25.4	16.7	24.0	7,735
RM-10	94.4	5.60	0	0	0	15,931	33.9	25.4	16.7	24.0	10,998
RM-11	94.4	5.60	0	0	0	12,677	38.7	29.3	20.2	11.8	8,041
RM-12	94.4	5.60	0	0	0	2,928	33.9	25.4	16.7	24.0	2,021
RM-13	94.4	5.60	0	0	0	17,051	43.0	27.4	14.0	15.6	10,738
RM-14	94.4	5.60	0	0	0	27,498	33.9	25.4	16.7	24.0	18,983
RM-15	94.4	5.60	0	0	0	46,611	33.5	31.6	22.1	12.8	41,135

Table C-4 (continued).

Site	Natural Gas					Natural Gas, EUR (Mg)	Natural Gas Liquids				Natural Gas Liquids, EUR (Mg)
	% C ₁	% C ₂	% C ₃	% C ₄	% C ₅₊		% C ₂	% C ₃	% C ₄	% C ₅₊	
RM-16	94.4	5.60	0	0	0	2,192	33.9	25.4	16.7	24.0	1,513
RM-17	94.4	5.60	0	0	0	103,060	37.0	32.9	19.2	11.0	84,703
RM-18	94.4	5.60	0	0	0	13,386	33.9	25.4	16.7	24.0	9,241
RM-19	94.4	5.60	0	0	0	28,106	34.6	25.6	17.9	22.0	17,240
Weighted Average	97.3	2.61	0.120	0.007	0.003		38.6	27.8	18.8	14.8	

Table C-5. fraction of production by salable product (mass basis)

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30-yr well lifetime)	Natural Gas, fraction of total production (mass basis)	Natural Gas Liquids, fraction of total production (mass basis)	Oil, fraction of total production (mass basis)
AP-1	59,005	0	1	0	0
AP-2	211,289	0	1	0	0
AP-3	576,497	0	1	0	0
AP-4	836,980	0	1	0	0
AP-5	136,625	0	1	0	0
AP-6	516,860	0	1	0	0
AP-7	565,897	0	1	0	0
AP-8	87,743	1,037	0.62	0.36	0.01
AP-9	455,027	0	1	0	0
AP-10	614,983	0	1	0	0
AP-11	113,272	0	1	0	0
AP-12	160,139	0	1	0	0
AP-13	599,499	0	1	0	0
AP-14	387,376	0	1	0	0
AP-15	184,176	0	1	0	0
AP-16	128,401	0	1	0	0
AP-17	176,088	0	1	0	0
AP-18	988,031	0	1	0	0
AP-19	410,082	0	1	0	0
AP-20	305,353	0	1	0	0
AP-21	147,975	0	1	0	0
AP-22	1,523,474	0	1	0	0
AP-23	529,064	0	1	0	0
AP-24	3,140,582	0	1	0	0
AP-25	217,440	0	1	0	0
AP-26	540,977	0	1	0	0
AP-27	281,558	0	1	0	0
AP-28	105,196	0	1	0	0
AP-29	773,016	0	1	0	0
AP-31	540,977	0	1	0	0
AP-32	605,927	0	1	0	0
AP-33	1,417,505	0	1	0	0
AP-34	975,788	0	1	0	0
AP-35	141,472	0	1	0	0
AP-36	62,444	0	1	0	0
AP-37	347,208	0	1	0	0

Table C-5 (continued).

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30-yr well lifetime)	Natural Gas, fraction of total production (mass basis)	Natural Gas Liquids, fraction of total production (mass basis)	Oil, fraction of total production (mass basis)
AP-38	53,663	0	1	0	0
AP-39	1,477,859	0	1	0	0
AP-40	220,702	0	0.98	0.02	0
AP-41	437,108	0	1	0	0
AP-42	1,121,524	0	1	0	0
AP-43	352,842	0	1	0	0
AP-45	209,670	0	1	0	0
AP-46	430,507	0	1	0	0
AP-47	640,901	0	1	0	0
GC-1	16,390	94,298	0.09	0.06	0.85
GC-2	356,441	511,946	0.22	0.19	0.59
GC-3	27,566	0	1	0	0
GC-4	855,109	455,392	0.37	0.29	0.35
GC-5	241,501	402,361	0.21	0.17	0.62
GC-6	403,970	197,954	0.44	0.23	0.33
GC-7	96,687	0	1	0	0
GC-8	84,653	0	1	0	0
GC-9	1,140,739	413,697	0.48	0.25	0.27
GC-10	1,495,472	0	1	0	0
GC-11	506,093	0	1	0	0
GC-12	1,105,696	0	1	0	0
GC-13	139,183	0	1	0	0
GC-14	480,788	400,389	0.25	0.29	0.45
GC-15	1,176,216	0	1	0	0
GC-16	221,711	796,204	0.14	0.08	0.78
GC-17	1,028,161	0	1	0	0
GC-18	1,018,236	0	1	0	0
GC-19	938,788	0	1	0	0
GC-20	64,113	0	1	0	0
GC-21	86,101	0	1	0	0
GC-22	65,716	0	1	0	0
GC-23	433,391	0	1	0	0
GC-24	286,622	234,422	0.29	0.26	0.45
GC-25	7,203	0	0.94	0.06	0
GC-26	1,174,402	221,271	0.56	0.28	0.16
GC-27	24,168	0	0.96	0.04	0

Table C-5 (continued).

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30-yr well lifetime)	Natural Gas, fraction of total production (mass basis)	Natural Gas Liquids, fraction of total production (mass basis)	Oil, fraction of total production (mass basis)
GC-28	782,134	0	1	0	0
GC-29	50,244	0	1	0	0
GC-30	321,101	0	1	0	0
GC-31	357,985	530,696	0.28	0.12	0.60
GC-32	386,840	0	1	0	0
GC-33	103,660	0	1	0	0
GC-34	1,265,190	0	1	0	0
GC-35	323,297	0	1	0	0
GC-36	22,905	0	1	0	0
GC-37	581,859	506,710	0.27	0.26	0.47
GC-38	119,358	0	1	0	0
GC-39	106,442	0	1	0	0
GC-40	16,306	0	1	0	0
GC-41	7,027	0	1	0	0
GC-42	44,292	0	1	0	0
GC-43	22,216	0	1	0	0
GC-44	7,344	0	0.90	0.10	0
GC-45	5,638	0	0.86	0.14	0
GC-46	89,328	1,698	0.84	0.15	0.02
GC-47	15,016	0	0.92	0.08	0
GC-48	24,479	0	1	0	0
GC-49	13,474	0	0.92	0.08	0
GC-50	347,560	0	1	0	0
GC-51	18,222	0	0.92	0.08	0
GC-52	735,815	0	1	0	0
GC-53	63,836	0	1	0	0
GC-54	146,328	247,524	0.27	0.10	0.63
GC-55	2,273,091	0	1	0	0
GC-56	107,109	0	1	0	0
GC-57	868,706	0	1	0	0
GC-58	174,867	0	1	0	0
MC-1	55,705	16,021	0.46	0.31	0.22
MC-4	256,972	0	0.97	0.03	0
MC-5	811	0	0.65	0.35	0
MC-6	94,923	11,371	0.63	0.26	0.11
MC-7	10,182	0	1	0	0

Table C-5 (continued).

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30-yr well lifetime)	Natural Gas, fraction of total production (mass basis)	Natural Gas Liquids, fraction of total production (mass basis)	Oil, fraction of total production (mass basis)
MC-8	90,266	0	1	0	0
MC-9	21,572	0	1	0	0
MC-10	162,090	0	1	0	0
MC-11	123,083	0	1	0	0
MC-12	25,449	17,387	0.35	0.24	0.41
MC-13	54,485	0	1	0	0
MC-14	15,015	0	0.68	0.32	0
MC-16	60,746	12,271	0.58	0.25	0.17
MC-17	316,873	0	1	0	0
MC-18	83,731	0	1	0	0
MC-19	175,342	0	1	0	0
MC-20	115,581	13,888	0.62	0.27	0.11
MC-21	35,790	43,603	0.31	0.14	0.55
MC-22	432,029	0	1	0	0
MC-23	535,876	0	1	0	0
MC-24	36,863	0	1	0	0
MC-25	132,448	0	1	0	0
MC-26	32,519	0	1	0	0
RM-2	32,012	8,346	0.47	0.32	0.21
RM-3	8,514	1,437	0.51	0.34	0.14
RM-4	28,973	23,968	0.34	0.21	0.45
RM-5	28,120	6,854	0.48	0.33	0.20
RM-6	12,352	1,453	0.53	0.37	0.11
RM-7	11,639	2,205	0.61	0.23	0.16
RM-8	14,976	14,023	0.31	0.21	0.48
RM-9	18,939	3,958	0.49	0.34	0.17
RM-10	26,929	8,271	0.45	0.31	0.23
RM-11	20,718	1,078	0.58	0.37	0.05
RM-12	4,950	2,999	0.37	0.25	0.38
RM-13	27,789	28,837	0.30	0.19	0.51
RM-14	46,480	7,367	0.51	0.35	0.14
RM-15	87,746	56,512	0.32	0.29	0.39
RM-16	3,705	975	0.47	0.32	0.21
RM-17	187,763	27,717	0.48	0.39	0.13
RM-18	22,627	7,387	0.45	0.31	0.25
RM-19	45,346	7,367	0.53	0.33	0.14

Table C-6. Allocation of total (sum of emissions from pneumatic controllers, pneumatic pumps, and equipment leaks) emissions (Mg) by product (mass basis) over a 30 year well lifetime

Site	Natural Gas	Natural Gas Liquids	Oil
AP-1	38.15	0	0
AP-2	41.17	0	0
AP-3	262.4	0	0
AP-4	90.22	0	0
AP-5	31.49	0	0
AP-6	66.60	0	0
AP-7	164.1	0	0
AP-8	105.0	61.47	1.968
AP-9	670.9	0	0
AP-10	674.5	0	0
AP-11	11.35	0	0
AP-12	119.4	0.4894	0
AP-13	298.5	0	0
AP-14	70.54	0	0
AP-15	0	0	0
AP-16	65.39	0	0
AP-17	353.3	0	0
AP-18	87.49	0	0
AP-19	116.0	0	0
AP-20	0	0	0
AP-21	153.2	0	0
AP-22	310.6	0	0
AP-23	320.6	0	0
AP-24	106.3	0	0
AP-25	1,758	0	0
AP-26	195.3	0	0
AP-27	848.0	0	0
AP-28	43.60	0	0
AP-29	163.8	0	0
AP-31	254.9	0	0
AP-32	98.70	0	0
AP-33	2,164	0	0
AP-34	2,184	0	0
AP-35	380.4	0	0
AP-36	9.082	0	0
AP-37	181.3	0	0

Table C-6 (continued).

Site	Natural Gas	Natural Gas Liquids	Oil
AP-38	494.7	0	0
AP-39	406.3	0	0
AP-40	63.19	1.598	0
AP-41	29.97	0	0
AP-42	1,052	0	0
AP-43	893.7	0	0
AP-45	42.38	0	0
AP-46	272.5	0	0
AP-47	42.69	0	0
GC-1	33.44	23.27	326.3
GC-2	138.9	123.9	377.5
GC-3	337.3	0	0
GC-4	259.8	204.0	247.0
GC-5	426.9	348.7	1,292
GC-6	407.7	215.6	305.5
GC-7	148.3	0	0
GC-8	75.38	0	0
GC-9	36.05	19.05	19.98
GC-10	847.7	0	0
GC-11	158.0	0	0
GC-12	755.7	0	0
GC-13	252.2	0	0
GC-14	215.8	246.7	385.2
GC-15	953.2	0	0
GC-16	230.8	133.6	1,309
GC-17	1,082	0	0
GC-18	298.5	0	0
GC-19	1,132	0	0
GC-20	72.66	0	0
GC-21	13.32	0	0
GC-22	347.2	0	0
GC-23	45.41	0	0
GC-24	192.7	172.0	298.3
GC-25	210.5	12.88	0
GC-26	514.6	253.2	144.7
GC-27	165.7	7.481	0
GC-28	213.2	0	0
GC-29	193.2	0	0

Table C-6 (continued).

Site	Natural Gas	Natural Gas Liquids	Oil
GC-30	479.2	0	0
GC-31	61.02	26.79	130.2
GC-32	294.3	0	0
GC-33	139.3	0	0
GC-34	845.6	0	0
GC-35	1,618	0	0
GC-36	607.9	0	0
GC-37	29.79	28.14	50.45
GC-38	119.3	0	0
GC-39	1,591	0	0
GC-40	180.7	0	0
GC-41	751.1	0	0
GC-42	534.6	0	0
GC-43	129.3	0	0
GC-44	3.826	0.4124	0
GC-45	91.92	14.94	0
GC-46	115.4	20.10	2.575
GC-47	267.5	23.16	0
GC-48	147.7	0	0
GC-49	265.8	23.02	0
GC-50	144.1	0	0
GC-51	297.2	24.62	0
GC-52	540.4	0	0
GC-53	455.9	0	0
GC-54	16.46	6.189	38.31
GC-55	594.0	0	0
GC-56	132.6	0	0
GC-57	227.7	0	0
GC-58	848.4	0	0
MC-1	252.9	172.2	122.3
MC-4	393.0	12.42	0
MC-5	184.6	98.19	0
MC-6	998.7	420.7	170.0
MC-7	62.67	0	0
MC-8	498.6	0	0
MC-9	459.3	0	0
MC-10	94.46	0	0
MC-11	871.0	0	0

Table C-6 (continued).

Site	Natural Gas	Natural Gas Liquids	Oil
MC-12	167.2	117.3	194.4
MC-13	1,161	0	0
MC-14	203.8	94.13	0
MC-16	308.4	130.8	88.73
MC-17	1,762	0	0
MC-18	181.3	0	0
MC-19	1,065	0	0
MC-20	259.0	115.2	44.96
MC-21	91.60	39.60	159.9
MC-22	4,140	0	0
MC-23	413.2	0	0
MC-24	164.1	0	0
MC-25	152.0	0	0
MC-26	110.2	0	0
RM-2	13.07	9.023	5.760
RM-3	17.70	11.83	4.985
RM-4	3.800	2.331	5.071
RM-5	724.3	500.0	298.5
RM-6	375.5	259.2	74.66
RM-7	392.8	147.8	102.4
RM-8	9.527	6.577	15.08
RM-9	4.148	2.864	1.465
RM-10	69.47	47.96	36.07
RM-11	0.8804	0.5584	0.0749
RM-12	2.119	1.463	2.170
RM-13	0.1823	0.1148	0.3084
RM-14	335.5	231.6	89.88
RM-15	0	0	0
RM-16	4.821	3.328	2.144
RM-17	157.3	129.2	42.29
RM-18	19.44	13.42	10.73
RM-19	44.23	27.13	11.59

Table C-7. Ratio of methane emissions (over a 30 year period) to estimated ultimate recovery of Natural Gas, Natural Gas Liquid, and Oil (Mass Basis) over a 30 year well lifetime.

Site	Mass Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
AP-1	0.0007	NA	NA
AP-2	0.0002	NA	NA
AP-3	0.0005	NA	NA
AP-4	0.0001	NA	NA
AP-5	0.0002	NA	NA
AP-6	0.0001	NA	NA
AP-7	0.0003	NA	NA
AP-8	0.0019	0.0019	0.0019
AP-9	0.0015	NA	NA
AP-10	0.0011	NA	NA
AP-11	0.0001	NA	NA
AP-12	0.0008	0.0007	NA
AP-13	0.0005	NA	NA
AP-14	0.0002	NA	NA
AP-15	0.0E+00	NA	NA
AP-16	0.0005	NA	NA
AP-17	0.0020	NA	NA
AP-18	9.1E-05	NA	NA
AP-19	0.0003	NA	NA
AP-20	0.0E+00	NA	NA
AP-21	0.0011	NA	NA
AP-22	0.0002	NA	NA
AP-23	0.0006	NA	NA
AP-24	3.5E-05	NA	NA
AP-25	0.0082	NA	NA
AP-26	0.0004	NA	NA
AP-27	0.0031	NA	NA
AP-28	0.0004	NA	NA
AP-29	0.0002	NA	NA
AP-31	0.0005	NA	NA
AP-32	0.0002	NA	NA
AP-33	0.0016	NA	NA
AP-34	0.0023	NA	NA
AP-35	0.0027	NA	NA

Table C-7 (continued).

Site	Mass Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
AP-36	0.0001	NA	NA
AP-37	0.0005	NA	NA
AP-38	0.0093	NA	NA
AP-39	0.0003	NA	NA
AP-40	0.0003	0.0003	NA
AP-41	7.0E-05	NA	NA
AP-42	0.0010	NA	NA
AP-43	0.0026	NA	NA
AP-45	0.0002	NA	NA
AP-46	0.0006	NA	NA
AP-47	6.8E-05	NA	NA
GC-1	0.0036	0.0035	0.0035
GC-2	0.0008	0.0007	0.0007
GC-3	0.0122	NA	NA
GC-4	0.0006	0.0005	0.0005
GC-5	0.0033	0.0032	0.0032
GC-6	0.0016	0.0015	0.0015
GC-7	0.0015	NA	NA
GC-8	0.0009	NA	NA
GC-9	5.0E-05	4.8E-05	4.8E-05
GC-10	0.0006	NA	NA
GC-11	0.0003	NA	NA
GC-12	0.0007	NA	NA
GC-13	0.0018	NA	NA
GC-14	0.0010	0.0010	0.0010
GC-15	0.0008	NA	NA
GC-16	0.0017	0.0016	0.0016
GC-17	0.0011	NA	NA
GC-18	0.0003	NA	NA
GC-19	0.0012	NA	NA
GC-20	0.0011	NA	NA
GC-21	0.0002	NA	NA
GC-22	0.0053	NA	NA
GC-23	0.0001	NA	NA
GC-24	0.0013	0.0013	0.0013
GC-25	0.0319	0.0310	NA

Table C-7 (continued).

Site	Mass Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
GC-26	0.0007	0.0007	0.0007
GC-27	0.0074	0.0072	NA
GC-28	0.0003	NA	NA
GC-29	0.0039	NA	NA
GC-30	0.0015	NA	NA
GC-31	0.0003	0.0002	0.0002
GC-32	0.0008	NA	NA
GC-33	0.0013	NA	NA
GC-34	0.0007	NA	NA
GC-35	0.0050	NA	NA
GC-36	0.0266	NA	NA
GC-37	0.0001	1.0E-04	1.0E-04
GC-38	0.0010	NA	NA
GC-39	0.0151	NA	NA
GC-40	0.0111	NA	NA
GC-41	0.1082	NA	NA
GC-42	0.0121	NA	NA
GC-43	0.0059	NA	NA
GC-44	0.0006	0.0006	NA
GC-45	0.0195	0.0190	NA
GC-46	0.0016	0.0015	0.0015
GC-47	0.0199	0.0194	NA
GC-48	0.0060	NA	NA
GC-49	0.0220	0.0214	NA
GC-50	0.0004	NA	NA
GC-51	0.0181	0.0177	NA
GC-52	0.0007	NA	NA
GC-53	0.0071	NA	NA
GC-54	0.0002	0.0002	0.0002
GC-55	0.0003	NA	NA
GC-56	0.0012	NA	NA
GC-57	0.0003	NA	NA
GC-58	0.0049	NA	NA
MC-1	0.0078	0.0076	0.0076
MC-4	0.0016	0.0016	NA
MC-5	0.3580	0.3487	NA

Table C-7 (continued).

Site	Mass Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
MC-6	0.0154	0.0150	0.0150
MC-7	0.0062	NA	NA
MC-8	0.0056	NA	NA
MC-9	0.0216	NA	NA
MC-10	0.0006	NA	NA
MC-11	0.0072	NA	NA
MC-12	0.0115	0.0112	0.0112
MC-13	0.0216	NA	NA
MC-14	0.0204	0.0198	NA
MC-16	0.0074	0.0072	0.0072
MC-17	0.0056	NA	NA
MC-18	0.0022	NA	NA
MC-19	0.0062	NA	NA
MC-20	0.0033	0.0032	0.0032
MC-21	0.0038	0.0037	0.0037
MC-22	0.0097	NA	NA
MC-23	0.0008	NA	NA
MC-24	0.0045	NA	NA
MC-25	0.0012	NA	NA
MC-26	0.0034	NA	NA
RM-2	0.0007	0.0007	0.0007
RM-3	0.0036	0.0035	0.0035
RM-4	0.0002	0.0002	0.0002
RM-5	0.0447	0.0435	0.0435
RM-6	0.0528	0.0514	0.0514
RM-7	0.0477	0.0464	0.0464
RM-8	0.0011	0.0011	0.0011
RM-9	0.0004	0.0004	0.0004
RM-10	0.0045	0.0044	0.0044
RM-11	7.1E-05	6.9E-05	6.9E-05
RM-12	0.0007	0.0007	0.0007
RM-13	1.1E-05	1.1E-05	1.1E-05
RM-14	0.0125	0.0122	0.0122
RM-15	0	0	0
RM-16	0.0023	0.0022	0.0022
RM-17	0.0016	0.0015	0.0015

Table C-7 (continued).

Site	Mass Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
RM-18	0.0015	0.0015	0.0015
RM-19	0.0016	0.0016	0.0016

C.5 Energy based allocation of emissions

Once production is split into each of the salable products (Natural Gas, Natural Gas Liquids, and Oil), as discussed in section C4, Estimated Ultimate Recovery (EUR) is converted from a mass base to an energy basis, using the conversion factors from section C1. Allocation of emissions is determined based on the heating value of each salable product.

For the case of the higher heating value of Natural Gas Liquids, a *weighted average heating value* is calculated for each site, based on its non-methane hydrocarbon gases composition, where the weighted average energy content is:

$$\langle Energy_{NMVOC} \rangle = \frac{1}{\text{non methane hydrocarbon}} (C_2 \times Energy_{C_2} + C_3 \times Energy_{C_3} + C_4 \times Energy_{C_4} + C_{5+} \times Energy_{C_{5+}})$$

(C5.1)

where $Energy_{C_n}$ refers to the heating value of the individual components from the gas.

Even though a fixed (natural gas) heating value is used for the conversion of salable natural gas in a mass base to an energy base (reported in Table C-1) for all sites, the average heating value of salable natural gas is calculated for each site to assess the potential impact of this assumption. Table C-8 shows the average heating value of both Natural Gas and Natural Gas Liquids. Table C-9 shows the fraction of production for each of the salable products (energy basis). Table C-10 shows the allocation of emissions from pneumatic controllers, pneumatic pumps, and equipment leaks over a 30 year period to each of the salable products, based. Table C-11 reports the ratio of allocated emissions to production for each of the three products: Natural Gas, Natural Gas Liquids, and Oil.

Table C-8. Average heating value of Natural Gas and Natural Gas Liquids for each site. For further calculations, in the case Natural Gas (Table C-1), a fixed higher heating value of 1,027 BTU/scf was assumed. In the case of Natural Gas Liquids, the average higher heating value at each site was used.

Site	Natural Gas, average higher heating value (BTU/scf)	Natural Gas Liquids, average higher heating value (BTU/scf)
AP-1	1,025	NA
AP-2	1,029	NA
AP-3	1,027	NA
AP-4	1,022	NA
AP-5	1,027	NA
AP-6	1,028	NA
AP-7	1,028	NA
AP-8	1,033	2,180
AP-9	1,022	NA
AP-10	1,022	NA
AP-11	1,025	NA
AP-12	1,032	2,567
AP-13	1,026	NA
AP-14	1,031	NA
AP-15	1,021	NA
AP-16	1,032	NA
AP-17	1,024	NA
AP-18	1,031	NA
AP-19	1,030	NA
AP-20	1,021	NA
AP-21	1,028	NA
AP-22	1,026	NA
AP-23	1,026	NA
AP-24	1,028	NA
AP-25	1,023	NA
AP-26	1,027	NA
AP-27	1,023	NA
AP-28	1,027	NA
AP-29	1,028	NA
AP-31	1,027	NA
AP-32	1,027	NA
AP-33	1,028	NA
AP-34	1,028	NA
AP-35	1,029	NA

Table C-8 (continued).

Site	Natural Gas, average heating value (BTU/scf)	Natural Gas Liquids, average heating value (BTU/scf)
AP-36	1,027	NA
AP-37	1,031	NA
AP-38	1,020	NA
AP-39	1,029	NA
AP-40	1,033	1,994
AP-41	1,027	NA
AP-42	1,029	NA
AP-43	1,031	NA
AP-45	1,028	NA
AP-46	1,026	NA
AP-47	1,031	NA
GC-1	1,033	2,320
GC-2	1,033	2,367
GC-3	1,010	NA
GC-4	1,033	2,341
GC-5	1,033	2,370
GC-6	1,033	2,262
GC-7	1,010	NA
GC-8	1,012	NA
GC-9	1,033	2,264
GC-10	1,010	NA
GC-11	1,011	NA
GC-12	1,010	NA
GC-13	1,011	NA
GC-14	1,033	2,490
GC-15	1,011	NA
GC-16	1,033	2,229
GC-17	1,010	NA
GC-18	1,010	NA
GC-19	1,011	NA
GC-20	1,010	NA
GC-21	1,010	NA
GC-22	1,011	NA
GC-23	1,010	NA
GC-24	1,033	2,408
GC-25	1,033	2,746
GC-26	1,033	2,314

Table C-8 (continued).

Site	Natural Gas, average heating value (BTU/scf)	Natural Gas Liquids, average heating value (BTU/scf)
GC-27	1,033	3,105
GC-28	1,011	NA
GC-29	1,011	NA
GC-30	1,010	NA
GC-31	1,033	2,277
GC-32	1,011	NA
GC-33	1,012	NA
GC-34	1,010	NA
GC-35	1,011	NA
GC-36	1,010	NA
GC-37	1,033	2,441
GC-38	1,010	NA
GC-39	1,020	NA
GC-40	1,011	NA
GC-41	1,020	NA
GC-42	1,010	NA
GC-43	1,020	NA
GC-44	1,033	3,327
GC-45	1,033	2,535
GC-46	1,033	2,430
GC-47	1,033	2,792
GC-48	1,011	NA
GC-49	1,033	2,792
GC-50	1,018	NA
GC-51	1,033	2,907
GC-52	1,011	NA
GC-53	1,010	NA
GC-54	1,033	2,241
GC-55	1,010	NA
GC-56	1,013	NA
GC-57	1,011	NA
GC-58	1,020	NA
MC-1	1,033	2,430
MC-4	1,033	2,047
MC-5	1,033	2,350
MC-6	1,033	2,388
MC-7	1,015	NA

Table C-8 (continued).

Site	Natural Gas, average heating value (BTU/scf)	Natural Gas Liquids, average heating value (BTU/scf)
MC-8	1,019	NA
MC-9	1,021	NA
MC-10	1,018	NA
MC-11	1,019	NA
MC-12	1,033	2,414
MC-13	1,021	NA
MC-14	1,033	2,324
MC-16	1,033	2,328
MC-17	1,019	NA
MC-18	1,022	NA
MC-19	1,022	NA
MC-20	1,033	2,322
MC-21	1,033	2,364
MC-22	1,021	NA
MC-23	1,017	NA
MC-24	1,019	NA
MC-25	1,017	NA
MC-26	1,019	NA
RM-2	1,033	2,447
RM-3	1,033	2,389
RM-4	1,033	2,430
RM-5	1,033	2,447
RM-6	1,033	2,447
RM-7	1,033	2,350
RM-8	1,033	2,447
RM-9	1,033	2,447
RM-10	1,033	2,447
RM-11	1,033	2,324
RM-12	1,033	2,447
RM-13	1,033	2,283
RM-14	1,033	2,447
RM-15	1,033	2,391
RM-16	1,033	2,447
RM-17	1,033	2,329
RM-18	1,033	2,447
RM-19	1,033	2,430

Table C-8 (continued).

Site	Natural Gas, average heating value (BTU/scf)	Natural Gas Liquids, average heating value (BTU/scf)
Weighted Average	1,022	2,349

Table C-9. Fraction of production by product (energy basis).

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30-yr well lifetime)	Natural Gas, fraction of total production energy basis)	Natural Gas Liquids, fraction of total production energy basis)	Oil, fraction of total production energy basis)
AP-1	59,005	0	1	0	0
AP-2	211,289	0	1	0	0
AP-3	576,497	0	1	0	0
AP-4	836,980	0	1	0	0
AP-5	136,625	0	1	0	0
AP-6	516,860	0	1	0	0
AP-7	565,897	0	1	0	0
AP-8	87,743	1,037	0.64	0.35	0.01
AP-9	455,027	0	1	0	0
AP-10	614,983	0	1	0	0
AP-11	113,272	0	1	0	0
AP-12	160,139	0	1.00	0.004	0
AP-13	599,499	0	1	0	0
AP-14	387,376	0	1	0	0
AP-15	184,176	0	1	0	0
AP-16	128,401	0	1	0	0
AP-17	176,088	0	1	0	0
AP-18	988,031	0	1	0	0
AP-19	410,082	0	1	0	0
AP-20	305,353	0	1	0	0
AP-21	147,975	0	1	0	0
AP-22	1,523,474	0	1	0	0
AP-23	529,064	0	1	0	0
AP-24	3,140,582	0	1	0	0
AP-25	217,440	0	1	0	0
AP-26	540,977	0	1	0	0
AP-27	281,558	0	1	0	0
AP-28	105,196	0	1	0	0
AP-29	773,016	0	1	0	0
AP-31	540,977	0	1	0	0
AP-32	605,927	0	1	0	0
AP-33	1,417,505	0	1	0	0
AP-34	975,788	0	1	0	0
AP-35	141,472	0	1	0	0

Table C-9 (continued).

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30-yr well lifetime)	Natural Gas, fraction of total production energy basis)	Natural Gas Liquids, fraction of total production energy basis)	Oil, fraction of total production energy basis)
AP-36	62,444	0	1	0	0
AP-37	347,208	0	1	0	0
AP-38	53,663	0	1	0	0
AP-39	1,477,859	0	1	0	0
AP-40	220,702	0	0.98	0.02	0
AP-41	437,108	0	1	0	0
AP-42	1,121,524	0	1	0	0
AP-43	352,842	0	1	0	0
AP-45	209,670	0	1	0	0
AP-46	430,507	0	1	0	0
AP-47	640,901	0	1	0	0
GC-1	16,390	94,298	0.10	0.06	0.84
GC-2	356,441	511,946	0.24	0.19	0.57
GC-3	27,566	0	1	0	0
GC-4	855,109	455,392	0.39	0.28	0.33
GC-5	241,501	402,361	0.23	0.17	0.61
GC-6	403,970	197,954	0.46	0.23	0.31
GC-7	96,687	0	1	0	0
GC-8	84,653	0	1	0	0
GC-9	1,140,739	413,697	0.51	0.25	0.25
GC-10	1,495,472	0	1	0	0
GC-11	506,093	0	1	0	0
GC-12	1,105,696	0	1	0	0
GC-13	139,183	0	1	0	0
GC-14	480,788	400,389	0.28	0.29	0.44
GC-15	1,176,216	0	1	0	0
GC-16	221,711	796,204	0.15	0.08	0.77
GC-17	1,028,161	0	1	0	0
GC-18	1,018,236	0	1	0	0
GC-19	938,788	0	1	0	0
GC-20	64,113	0	1	0	0
GC-21	86,101	0	1	0	0
GC-22	65,716	0	1	0	0
GC-23	433,391	0	1	0	0

Table C-9 (continued).

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30-yr well lifetime)	Natural Gas, fraction of total production energy basis)	Natural Gas Liquids, fraction of total production energy basis)	Oil, fraction of total production energy basis)
GC-24	286,622	234,422	0.31	0.26	0.43
GC-25	7,203	0	0.95	0.05	0
GC-26	1,174,402	221,271	0.59	0.26	0.15
GC-27	24,168	0	0.96	0.04	0
GC-28	782,134	0	1	0	0
GC-29	50,244	0	1	0	0
GC-30	321,101	0	1	0	0
GC-31	357,985	530,696	0.30	0.12	0.57
GC-32	386,840	0	1	0	0
GC-33	103,660	0	1	0	0
GC-34	1,265,190	0	1	0	0
GC-35	323,297	0	1	0	0
GC-36	22,905	0	1	0	0
GC-37	581,859	506,710	0.30	0.26	0.45
GC-38	119,358	0	1	0	0
GC-39	106,442	0	1	0	0
GC-40	16,306	0	1	0	0
GC-41	7,027	0	1	0	0
GC-42	44,292	0	1	0	0
GC-43	22,216	0	1	0	0
GC-44	7,344	0	0.91	0.09	0
GC-45	5,638	0	0.87	0.13	0
GC-46	89,328	1,698	0.85	0.13	0.02
GC-47	15,016	0	0.93	0.07	0
GC-48	24,479	0	1	0	0
GC-49	13,474	0	0.93	0.07	0
GC-50	347,560	0	1	0	0
GC-51	18,222	0	0.93	0.07	0
GC-52	735,815	0	1	0	0
GC-53	63,836	0	1	0	0
GC-54	146,328	247,524	0.29	0.10	0.61
GC-55	2,273,091	0	1	0	0
GC-56	107,109	0	1	0	0
GC-57	868,706	0	1	0	0

Table C-9 (continued).

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30-yr well lifetime)	Natural Gas, fraction of total production energy basis)	Natural Gas Liquids, fraction of total production energy basis)	Oil, fraction of total production energy basis)
GC-58	174,867	0	1	0	0
MC-1	55,705	16,021	0.49	0.31	0.20
MC-4	256,972	0	0.97	0.03	0
MC-5	811	0	0.67	0.33	0
MC-6	94,923	11,371	0.65	0.25	0.10
MC-7	10,182	0	1	0	0
MC-8	90,266	0	1	0	0
MC-9	21,572	0	1	0	0
MC-10	162,090	0	1	0	0
MC-11	123,083	0	1	0	0
MC-12	25,449	17,387	0.38	0.24	0.38
MC-13	54,485	0	1	0	0
MC-14	15,015	0	0.70	0.30	0
MC-16	60,746	12,271	0.61	0.24	0.15
MC-17	316,873	0	1	0	0
MC-18	83,731	0	1	0	0
MC-19	175,342	0	1	0	0
MC-20	115,581	13,888	0.64	0.26	0.10
MC-21	35,790	43,603	0.35	0.14	0.52
MC-22	432,029	0	1	0	0
MC-23	535,876	0	1	0	0
MC-24	36,863	0	1	0	0
MC-25	132,448	0	1	0	0
MC-26	32,519	0	1	0	0
RM-2	32,012	8,346	0.49	0.31	0.20
RM-3	8,514	1,437	0.54	0.33	0.14
RM-4	28,973	23,968	0.36	0.20	0.43
RM-5	28,120	6,854	0.50	0.31	0.19
RM-6	12,352	1,453	0.55	0.35	0.10
RM-7	11,639	2,205	0.63	0.22	0.15
RM-8	14,976	14,023	0.33	0.21	0.47
RM-9	18,939	3,958	0.51	0.32	0.16
RM-10	26,929	8,271	0.48	0.30	0.22
RM-11	20,718	1,078	0.60	0.35	0.05

Table C-9 (continued).

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30-yr well lifetime)	Natural Gas, fraction of total production energy basis)	Natural Gas Liquids, fraction of total production energy basis)	Oil, fraction of total production energy basis)
RM-12	4,950	2,999	0.39	0.25	0.36
RM-13	27,789	28,837	0.32	0.19	0.49
RM-14	46,480	7,367	0.54	0.34	0.13
RM-15	87,746	56,512	0.35	0.28	0.38
RM-16	3,705	975	0.49	0.31	0.20
RM-17	187,763	27,717	0.50	0.38	0.12
RM-18	22,627	7,387	0.47	0.29	0.23
RM-19	45,346	7,367	0.56	0.31	0.13

Table C-10. Allocation of total (sum of emissions from pneumatic controllers, pneumatic pumps, and equipment leaks) emissions (Mg) by product (energy basis) over a 30 year well lifetime

Site	Natural Gas	Natural Gas Liquids	Oil
AP-1	38.15	0	0
AP-2	41.17	0	0
AP-3	262.4	0	0
AP-4	90.22	0	0
AP-5	31.49	0	0
AP-6	66.60	0	0
AP-7	164.1	0	0
AP-8	108.2	58.44	1.801
AP-9	670.9	0	0
AP-10	674.5	0	0
AP-11	11.35	0	0
AP-12	119.4	0.4491	0
AP-13	298.5	0	0
AP-14	70.54	0	0
AP-15	0	0	0
AP-16	65.39	0	0
AP-17	353.3	0	0
AP-18	87.49	0	0
AP-19	116.0	0	0
AP-20	0	0	0
AP-21	153.2	0	0
AP-22	310.6	0	0
AP-23	320.6	0	0
AP-24	106.3	0	0
AP-25	1,758	0	0
AP-26	195.3	0	0
AP-27	848.0	0	0
AP-28	43.60	0	0
AP-29	163.8	0	0
AP-31	254.9	0	0
AP-32	98.70	0	0
AP-33	2,164	0	0
AP-34	2,184	0	0
AP-35	380.4	0	0
AP-36	9.082	0	0
AP-37	181.3	0	0
AP-38	494.7	0	0

Table C-10 (continued).

Site	Natural Gas	Natural Gas Liquids	Oil
AP-39	406.3	0	0
AP-40	63.29	1.495	0
AP-41	29.97	0	0
AP-42	1,052	0	0
AP-43	893.7	0	0
AP-45	42.38	0	0
AP-46	272.5	0	0
AP-47	42.69	0	0
GC-1	37.18	23.68	322.1
GC-2	151.4	123.4	365.5
GC-3	337.3	0	0
GC-4	277.3	199.3	234.2
GC-5	466.3	347.7	1,254
GC-6	431.8	209.7	287.3
GC-7	148.3	0	0
GC-8	75.38	0	0
GC-9	37.97	18.42	18.69
GC-10	847.7	0	0
GC-11	158.0	0	0
GC-12	755.7	0	0
GC-13	252.2	0	0
GC-14	234.2	242.2	371.2
GC-15	953.2	0	0
GC-16	254.7	135.8	1,283
GC-17	1,082	0	0
GC-18	298.5	0	0
GC-19	1,132	0	0
GC-20	72.66	0	0
GC-21	13.32	0	0
GC-22	347.2	0	0
GC-23	45.41	0	0
GC-24	208.0	169.1	285.9
GC-25	211.8	11.60	0
GC-26	536.8	241.7	134.0
GC-27	166.5	6.650	0
GC-28	213.2	0	0
GC-29	193.2	0	0
GC-30	479.2	0	0
GC-31	66.11	26.62	125.2

Table C-10 (continued).

Site	Natural Gas	Natural Gas Liquids	Oil
GC-32	294.3	0	0
GC-33	139.3	0	0
GC-34	845.6	0	0
GC-35	1,618	0	0
GC-36	607.9	0	0
GC-37	32.24	27.65	48.49
GC-38	119.3	0	0
GC-39	1,591	0	0
GC-40	180.7	0	0
GC-41	751.1	0	0
GC-42	534.6	0	0
GC-43	129.3	0	0
GC-44	3.874	0.3641	0
GC-45	93.22	13.65	0
GC-46	117.2	18.48	2.324
GC-47	269.8	20.85	0
GC-48	147.7	0	0
GC-49	268.1	20.72	0
GC-50	144.1	0	0
GC-51	299.8	22.04	0
GC-52	540.4	0	0
GC-53	455.9	0	0
GC-54	17.86	6.172	36.92
GC-55	594.0	0	0
GC-56	132.6	0	0
GC-57	227.7	0	0
GC-58	848.4	0	0
MC-1	269.0	167.1	111.2
MC-4	393.8	11.59	0
MC-5	190.3	92.47	0
MC-6	1,039	399.3	151.2
MC-7	62.67	0	0
MC-8	498.6	0	0
MC-9	459.3	0	0
MC-10	94.46	0	0
MC-11	871.0	0	0
MC-12	181.8	116.4	180.7
MC-13	1,161	0	0
MC-14	209.3	88.55	0

Table C-10 (continued).

Site	Natural Gas	Natural Gas Liquids	Oil
MC-16	323.1	125.4	79.48
MC-17	1,762	0	0
MC-18	181.3	0	0
MC-19	1,065	0	0
MC-20	269.4	109.8	39.99
MC-21	100.8	39.82	150.4
MC-22	4,140	0	0
MC-23	413.2	0	0
MC-24	164.1	0	0
MC-25	152.0	0	0
MC-26	110.2	0	0
RM-2	13.78	8.620	5.451
RM-3	18.53	11.30	4.684
RM-4	4.067	2.264	4.871
RM-5	763.2	477.4	282.2
RM-6	393.2	245.9	70.17
RM-7	407.9	139.6	95.44
RM-8	10.24	6.403	14.54
RM-9	4.364	2.730	1.383
RM-10	73.39	45.90	34.19
RM-11	0.9134	0.5306	0.0697
RM-12	2.261	1.414	2.078
RM-13	0.1957	0.1128	0.2970
RM-14	352.1	220.2	84.65
RM-15	0	0	0
RM-16	5.084	3.180	2.029
RM-17	164.9	124.1	39.79
RM-18	20.56	12.86	10.18
RM-19	46.29	25.77	10.89

Table C-11. Ratio of methane emissions (over a 30 year period) to estimated ultimate recovery of Natural Gas, Natural Gas Liquid, and Oil (energy Basis) over a 30 year well lifetime.

Site	Energy Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
AP-1	0.0007	NA	NA
AP-2	0.0002	NA	NA
AP-3	0.0005	NA	NA
AP-4	0.0001	NA	NA
AP-5	0.0002	NA	NA
AP-6	0.0001	NA	NA
AP-7	0.0003	NA	NA
AP-8	0.0020	0.0018	0.0017
AP-9	0.0015	NA	NA
AP-10	0.0011	NA	NA
AP-11	0.0001	NA	NA
AP-12	0.0008	0.0007	NA
AP-13	0.0005	NA	NA
AP-14	0.0002	NA	NA
AP-15	0.0E+00	NA	NA
AP-16	0.0005	NA	NA
AP-17	0.0020	NA	NA
AP-18	9.1E-05	NA	NA
AP-19	0.0003	NA	NA
AP-20	0.0E+00	NA	NA
AP-21	0.0011	NA	NA
AP-22	0.0002	NA	NA
AP-23	0.0006	NA	NA
AP-24	3.5E-05	NA	NA
AP-25	0.0082	NA	NA
AP-26	0.0004	NA	NA
AP-27	0.0031	NA	NA
AP-28	0.0004	NA	NA
AP-29	0.0002	NA	NA
AP-31	0.0005	NA	NA
AP-32	0.0002	NA	NA
AP-33	0.0016	NA	NA
AP-34	0.0023	NA	NA
AP-35	0.0027	NA	NA

Table C-11 (continued).

Site	Energy Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
AP-36	0.0001	NA	NA
AP-37	0.0005	NA	NA
AP-38	0.0093	NA	NA
AP-39	0.0003	NA	NA
AP-40	0.0003	0.0003	NA
AP-41	7.0E-05	NA	NA
AP-42	0.0010	NA	NA
AP-43	0.0026	NA	NA
AP-45	0.0002	NA	NA
AP-46	0.0006	NA	NA
AP-47	6.8E-05	NA	NA
GC-1	0.0039	0.0035	0.0034
GC-2	0.0008	0.0007	0.0007
GC-3	0.0122	NA	NA
GC-4	0.0006	0.0005	0.0005
GC-5	0.0036	0.0032	0.0031
GC-6	0.0017	0.0015	0.0015
GC-7	0.0015	NA	NA
GC-8	0.0009	NA	NA
GC-9	5.2E-05	4.7E-05	4.5E-05
GC-10	0.0006	NA	NA
GC-11	0.0003	NA	NA
GC-12	0.0007	NA	NA
GC-13	0.0018	NA	NA
GC-14	0.0011	0.0009	0.0009
GC-15	0.0008	NA	NA
GC-16	0.0019	0.0017	0.0016
GC-17	0.0011	NA	NA
GC-18	0.0003	NA	NA
GC-19	0.0012	NA	NA
GC-20	0.0011	NA	NA
GC-21	0.0002	NA	NA
GC-22	0.0053	NA	NA
GC-23	0.0001	NA	NA
GC-24	0.0014	0.0013	0.0012
GC-25	0.0320	0.0279	NA

Table C-11 (continued).

Site	Energy Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
GC-26	0.0007	0.0006	0.0006
GC-27	0.0074	0.0064	NA
GC-28	0.0003	NA	NA
GC-29	0.0039	NA	NA
GC-30	0.0015	NA	NA
GC-31	0.0003	0.0002	0.0002
GC-32	0.0008	NA	NA
GC-33	0.0013	NA	NA
GC-34	0.0007	NA	NA
GC-35	0.0050	NA	NA
GC-36	0.0266	NA	NA
GC-37	0.0001	9.8E-05	9.6E-05
GC-38	0.0010	NA	NA
GC-39	0.0151	NA	NA
GC-40	0.0111	NA	NA
GC-41	0.1082	NA	NA
GC-42	0.0121	NA	NA
GC-43	0.0059	NA	NA
GC-44	0.0006	0.0005	NA
GC-45	0.0197	0.0173	NA
GC-46	0.0016	0.0014	0.0014
GC-47	0.0200	0.0174	NA
GC-48	0.0060	NA	NA
GC-49	0.0222	0.0193	NA
GC-50	0.0004	NA	NA
GC-51	0.0183	0.0158	NA
GC-52	0.0007	NA	NA
GC-53	0.0071	NA	NA
GC-54	0.0002	0.0002	0.0001
GC-55	0.0003	NA	NA
GC-56	0.0012	NA	NA
GC-57	0.0003	NA	NA
GC-58	0.0049	NA	NA
MC-1	0.0083	0.0074	0.0069
MC-4	0.0016	0.0015	NA
MC-5	0.3691	0.3284	NA

Table C-11 (continued).

Site	Energy Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
MC-6	0.0160	0.0142	0.0133
MC-7	0.0062	NA	NA
MC-8	0.0056	NA	NA
MC-9	0.0216	NA	NA
MC-10	0.0006	NA	NA
MC-11	0.0072	NA	NA
MC-12	0.0125	0.0111	0.0104
MC-13	0.0216	NA	NA
MC-14	0.0209	0.0187	NA
MC-16	0.0078	0.0069	0.0065
MC-17	0.0056	NA	NA
MC-18	0.0022	NA	NA
MC-19	0.0062	NA	NA
MC-20	0.0035	0.0031	0.0029
MC-21	0.0041	0.0037	0.0034
MC-22	0.0097	NA	NA
MC-23	0.0008	NA	NA
MC-24	0.0045	NA	NA
MC-25	0.0012	NA	NA
MC-26	0.0034	NA	NA
RM-2	0.0007	0.0007	0.0007
RM-3	0.0037	0.0033	0.0033
RM-4	0.0002	0.0002	0.0002
RM-5	0.0471	0.0416	0.0412
RM-6	0.0553	0.0488	0.0483
RM-7	0.0495	0.0439	0.0433
RM-8	0.0012	0.0010	0.0010
RM-9	0.0004	0.0004	0.0003
RM-10	0.0047	0.0042	0.0041
RM-11	7.4E-05	6.6E-05	6.5E-05
RM-12	0.0008	0.0007	0.0007
RM-13	1.2E-05	1.1E-05	1.0E-05
RM-14	0.0131	0.0116	0.0115
RM-15	0	0	0
RM-16	0.0024	0.0021	0.0021
RM-17	0.0016	0.0015	0.0014

Table C-11 (continued).

Site	Energy Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
RM-18	0.0016	0.0014	0.0014
RM-19	0.0017	0.0015	0.0015

C.6 Price based allocation of emissions

Once production is split into each of the salable products (Natural Gas, Natural Gas Liquids, and Oil), as discussed in section C4, Estimated Ultimate Recovery is converted from a mass basis into a price base, using the conversion factors from section C1. Allocation of emissions is determined based on the price of each salable product.

Table C-12 shows the fraction of production for each of the salable products (price basis). Table C-13 shows the allocation of emissions from pneumatic controllers, pneumatic pumps, and equipment leaks over a 30 year period to each of the salable products, based. Table C-14 reports the ratio of allocated emissions to production for each of the three products: Natural Gas, Natural Gas Liquids, and Oil.

Table C-12. Fraction of production by product (price basis).

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30- yr well lifetime)	Natural Gas, fraction of total production price basis)	Natural Gas Liquids, fraction of total production price basis)	Oil, fraction of total production price basis)
AP-1	59,005	0	1	0	0
AP-2	211,289	0	1	0	0
AP-3	576,497	0	1	0	0
AP-4	836,980	0	1	0	0
AP-5	136,625	0	1	0	0
AP-6	516,860	0	1	0	0
AP-7	565,897	0	1	0	0
AP-8	87,743	1,037	0.38	0.59	0.03
AP-9	455,027	0	1	0	0
AP-10	614,983	0	1	0	0
AP-11	113,272	0	1	0	0
AP-12	160,139	0	0.99	0.01	0
AP-13	599,499	0	1	0	0
AP-14	387,376	0	1	0	0
AP-15	184,176	0	1	0	0
AP-16	128,401	0	1	0	0
AP-17	176,088	0	1	0	0
AP-18	988,031	0	1	0	0
AP-19	410,082	0	1	0	0
AP-20	305,353	0	1	0	0
AP-21	147,975	0	1	0	0
AP-22	1,523,474	0	1	0	0
AP-23	529,064	0	1	0	0
AP-24	3,140,582	0	1	0	0
AP-25	217,440	0	1	0	0
AP-26	540,977	0	1	0	0
AP-27	281,558	0	1	0	0
AP-28	105,196	0	1	0	0
AP-29	773,016	0	1	0	0
AP-31	540,977	0	1	0	0
AP-32	605,927	0	1	0	0
AP-33	1,417,505	0	1	0	0
AP-34	975,788	0	1	0	0
AP-35	141,472	0	1	0	0
AP-36	62,444	0	1	0	0
AP-37	347,208	0	1	0	0

Table C-12 (continued).

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30- yr well lifetime)	Natural Gas, fraction of total production price basis)	Natural Gas Liquids, fraction of total production price basis)	Oil, fraction of total production price basis)
AP-38	53,663	0	1	0	0
AP-39	1,477,859	0	1	0	0
AP-40	220,702	0	0.94	0.06	0
AP-41	437,108	0	1	0	0
AP-42	1,121,524	0	1	0	0
AP-43	352,842	0	1	0	0
AP-45	209,670	0	1	0	0
AP-46	430,507	0	1	0	0
AP-47	640,901	0	1	0	0
GC-1	16,390	94,298	0.02	0.04	0.93
GC-2	356,441	511,946	0.07	0.16	0.76
GC-3	27,566	0	1	0	0
GC-4	855,109	455,392	0.15	0.30	0.55
GC-5	241,501	402,361	0.07	0.14	0.79
GC-6	403,970	197,954	0.19	0.26	0.55
GC-7	96,687	0	1	0	0
GC-8	84,653	0	1	0	0
GC-9	1,140,739	413,697	0.22	0.30	0.48
GC-10	1,495,472	0	1	0	0
GC-11	506,093	0	1	0	0
GC-12	1,105,696	0	1	0	0
GC-13	139,183	0	1	0	0
GC-14	480,788	400,389	0.09	0.27	0.64
GC-15	1,176,216	0	1	0	0
GC-16	221,711	796,204	0.04	0.06	0.90
GC-17	1,028,161	0	1	0	0
GC-18	1,018,236	0	1	0	0
GC-19	938,788	0	1	0	0
GC-20	64,113	0	1	0	0
GC-21	86,101	0	1	0	0
GC-22	65,716	0	1	0	0
GC-23	433,391	0	1	0	0
GC-24	286,622	234,422	0.11	0.24	0.65
GC-25	7,203	0	0.87	0.13	0
GC-26	1,174,402	221,271	0.30	0.38	0.33
GC-27	24,168	0	0.90	0.10	0

Table C-12 (continued).

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30- yr well lifetime)	Natural Gas, fraction of total production price basis)	Natural Gas Liquids, fraction of total production price basis)	Oil, fraction of total production price basis)
GC-28	782,134	0	1	0	0
GC-29	50,244	0	1	0	0
GC-30	321,101	0	1	0	0
GC-31	357,985	530,696	0.10	0.11	0.80
GC-32	386,840	0	1	0	0
GC-33	103,660	0	1	0	0
GC-34	1,265,190	0	1	0	0
GC-35	323,297	0	1	0	0
GC-36	22,905	0	1	0	0
GC-37	581,859	506,710	0.10	0.24	0.66
GC-38	119,358	0	1	0	0
GC-39	106,442	0	1	0	0
GC-40	16,306	0	1	0	0
GC-41	7,027	0	1	0	0
GC-42	44,292	0	1	0	0
GC-43	22,216	0	1	0	0
GC-44	7,344	0	0.79	0.21	0
GC-45	5,638	0	0.71	0.29	0
GC-46	89,328	1,698	0.65	0.29	0.06
GC-47	15,016	0	0.82	0.18	0
GC-48	24,479	0	1	0	0
GC-49	13,474	0	0.82	0.18	0
GC-50	347,560	0	1	0	0
GC-51	18,222	0	0.83	0.17	0
GC-52	735,815	0	1	0	0
GC-53	63,836	0	1	0	0
GC-54	146,328	247,524	0.09	0.09	0.82
GC-55	2,273,091	0	1	0	0
GC-56	107,109	0	1	0	0
GC-57	868,706	0	1	0	0
GC-58	174,867	0	1	0	0
MC-1	55,705	16,021	0.22	0.38	0.40
MC-4	256,972	0	0.92	0.08	0
MC-5	811	0	0.42	0.58	0
MC-6	94,923	11,371	0.37	0.40	0.24
MC-7	10,182	0	1	0	0

Table C-12 (continued).

Site	Gas EUR (Mg) (30-yr well lifetime)	Oil EUR (Mg) (30- yr well lifetime)	Natural Gas, fraction of total production price basis)	Natural Gas Liquids, fraction of total production price basis)	Oil, fraction of total production price basis)
MC-8	90,266	0	1	0	0
MC-9	21,572	0	1	0	0
MC-10	162,090	0	1	0	0
MC-11	123,083	0	1	0	0
MC-12	25,449	17,387	0.14	0.25	0.61
MC-13	54,485	0	1	0	0
MC-14	15,015	0	0.46	0.54	0
MC-16	60,746	12,271	0.31	0.34	0.34
MC-17	316,873	0	1	0	0
MC-18	83,731	0	1	0	0
MC-19	175,342	0	1	0	0
MC-20	115,581	13,888	0.36	0.41	0.23
MC-21	35,790	43,603	0.11	0.13	0.76
MC-22	432,029	0	1	0	0
MC-23	535,876	0	1	0	0
MC-24	36,863	0	1	0	0
MC-25	132,448	0	1	0	0
MC-26	32,519	0	1	0	0
RM-2	32,012	8,346	0.22	0.39	0.39
RM-3	8,514	1,437	0.26	0.45	0.29
RM-4	28,973	23,968	0.13	0.20	0.67
RM-5	28,120	6,854	0.23	0.40	0.37
RM-6	12,352	1,453	0.28	0.50	0.22
RM-7	11,639	2,205	0.33	0.32	0.34
RM-8	14,976	14,023	0.11	0.20	0.69
RM-9	18,939	3,958	0.24	0.42	0.34
RM-10	26,929	8,271	0.21	0.37	0.43
RM-11	20,718	1,078	0.34	0.55	0.11
RM-12	4,950	2,999	0.15	0.26	0.60
RM-13	27,789	28,837	0.11	0.17	0.72
RM-14	46,480	7,367	0.26	0.46	0.28
RM-15	87,746	56,512	0.12	0.28	0.60
RM-16	3,705	975	0.22	0.39	0.39
RM-17	187,763	27,717	0.24	0.51	0.25
RM-18	22,627	7,387	0.20	0.36	0.44
RM-19	45,346	7,367	0.28	0.44	0.29

Table C-13. Allocation of total (sum of emissions from pneumatic controllers, pneumatic pumps, and equipment leaks) emissions (Mg) by product (price basis) over a 30 year well lifetime.

Site	Natural Gas	Natural Gas Liquids	Oil
AP-1	38.15	0	0
AP-2	41.17	0	0
AP-3	262.4	0	0
AP-4	90.22	0	0
AP-5	31.49	0	0
AP-6	66.60	0	0
AP-7	164.1	0	0
AP-8	64.84	98.86	4.775
AP-9	670.9	0	0
AP-10	674.5	0	0
AP-11	11.35	0	0
AP-12	118.6	1.260	0
AP-13	298.5	0	0
AP-14	70.54	0	0
AP-15	0	0	0
AP-16	65.39	0	0
AP-17	353.3	0	0
AP-18	87.49	0	0
AP-19	116.0	0	0
AP-20	0	0	0
AP-21	153.2	0	0
AP-22	310.6	0	0
AP-23	320.6	0	0
AP-24	106.3	0	0
AP-25	1,758	0	0
AP-26	195.3	0	0
AP-27	848.0	0	0
AP-28	43.60	0	0
AP-29	163.8	0	0
AP-31	254.9	0	0
AP-32	98.70	0	0
AP-33	2,164	0	0
AP-34	2,184	0	0
AP-35	380.4	0	0
AP-36	9.082	0	0
AP-37	181.3	0	0
AP-38	494.7	0	0

Table C-13 (continued).

Site	Natural Gas	Natural Gas Liquids	Oil
AP-39	406.3	0	0
AP-40	60.74	4.052	0
AP-41	29.97	0	0
AP-42	1,052	0	0
AP-43	893.7	0	0
AP-45	42.38	0	0
AP-46	272.5	0	0
AP-47	42.69	0	0
GC-1	9.305	16.74	356.9
GC-2	45.79	105.4	489.1
GC-3	337.3	0	0
GC-4	105.0	213.2	392.6
GC-5	137.8	290.2	1,640
GC-6	174.7	239.6	514.5
GC-7	148.3	0	0
GC-8	75.38	0	0
GC-9	16.51	22.61	35.97
GC-10	847.7	0	0
GC-11	158.0	0	0
GC-12	755.7	0	0
GC-13	252.2	0	0
GC-14	77.51	226.4	543.8
GC-15	953.2	0	0
GC-16	67.47	101.6	1,504
GC-17	1,082	0	0
GC-18	298.5	0	0
GC-19	1,132	0	0
GC-20	72.66	0	0
GC-21	13.32	0	0
GC-22	347.2	0	0
GC-23	45.41	0	0
GC-24	70.67	162.3	430.1
GC-25	193.5	29.93	0
GC-26	270.2	343.6	298.7
GC-27	155.6	17.55	0
GC-28	213.2	0	0
GC-29	193.2	0	0
GC-30	479.2	0	0
GC-31	20.71	23.56	173.7

Table C-13 (continued).

Site	Natural Gas	Natural Gas Liquids	Oil
GC-32	294.3	0	0
GC-33	139.3	0	0
GC-34	845.6	0	0
GC-35	1,618	0	0
GC-36	607.9	0	0
GC-37	10.75	26.05	71.59
GC-38	119.3	0	0
GC-39	1,591	0	0
GC-40	180.7	0	0
GC-41	751.1	0	0
GC-42	534.6	0	0
GC-43	129.3	0	0
GC-44	3.349	0.8891	0
GC-45	75.60	31.27	0
GC-46	90.06	40.09	7.903
GC-47	238.6	52.08	0
GC-48	147.7	0	0
GC-49	237.1	51.75	0
GC-50	144.1	0	0
GC-51	266.5	55.33	0
GC-52	540.4	0	0
GC-53	455.9	0	0
GC-54	5.478	5.347	50.13
GC-55	594.0	0	0
GC-56	132.6	0	0
GC-57	227.7	0	0
GC-58	848.4	0	0
MC-1	119.4	209.4	218.5
MC-4	374.3	31.11	0
MC-5	119.2	163.6	0
MC-6	582.2	632.0	375.2
MC-7	62.67	0	0
MC-8	498.6	0	0
MC-9	459.3	0	0
MC-10	94.46	0	0
MC-11	871.0	0	0
MC-12	66.44	120.1	292.4
MC-13	1,161	0	0
MC-14	135.7	162.2	0

Table C-13 (continued).

Site	Natural Gas	Natural Gas Liquids	Oil
MC-16	165.7	181.8	180.5
MC-17	1,762	0	0
MC-18	181.3	0	0
MC-19	1,065	0	0
MC-20	149.3	171.8	98.10
MC-21	33.37	37.23	220.5
MC-22	4,140	0	0
MC-23	413.2	0	0
MC-24	164.1	0	0
MC-25	152.0	0	0
MC-26	110.2	0	0
RM-2	6.166	10.89	10.80
RM-3	8.985	15.47	10.05
RM-4	1.423	2.236	7.543
RM-5	345.8	610.9	566.1
RM-6	199.5	352.3	157.6
RM-7	214.2	207.0	221.8
RM-8	3.444	6.084	21.66
RM-9	2.033	3.591	2.853
RM-10	31.79	56.15	65.56
RM-11	0.5082	0.8338	0.1717
RM-12	0.8416	1.487	3.424
RM-13	0.0648	0.1055	0.4353
RM-14	171.5	302.9	182.5
RM-15	0	0	0
RM-16	2.271	4.011	4.011
RM-17	78.37	166.7	83.73
RM-18	8.792	15.53	19.27
RM-19	22.96	36.09	23.90

Table C-14. Ratio of methane emissions (over a 30 year period) to estimated ultimate recovery of Natural Gas, Natural Gas Liquid, and Oil (price Basis) over a 30 year well lifetime:

Site	Price Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
AP-1	0.0007	NA	NA
AP-2	0.0002	NA	NA
AP-3	0.0005	NA	NA
AP-4	0.0001	NA	NA
AP-5	0.0002	NA	NA
AP-6	0.0001	NA	NA
AP-7	0.0003	NA	NA
AP-8	0.0012	0.0031	0.0046
AP-9	0.0015	NA	NA
AP-10	0.0011	NA	NA
AP-11	0.0001	NA	NA
AP-12	0.0008	0.0019	NA
AP-13	0.0005	NA	NA
AP-14	0.0002	NA	NA
AP-15	0.0E+00	NA	NA
AP-16	0.0005	NA	NA
AP-17	0.0020	NA	NA
AP-18	9.1E-05	NA	NA
AP-19	0.0003	NA	NA
AP-20	0.0E+00	NA	NA
AP-21	0.0011	NA	NA
AP-22	0.0002	NA	NA
AP-23	0.0006	NA	NA
AP-24	3.5E-05	NA	NA
AP-25	0.0082	NA	NA
AP-26	0.0004	NA	NA
AP-27	0.0031	NA	NA
AP-28	0.0004	NA	NA
AP-29	0.0002	NA	NA
AP-31	0.0005	NA	NA
AP-32	0.0002	NA	NA
AP-33	0.0016	NA	NA
AP-34	0.0023	NA	NA
AP-35	0.0027	NA	NA

Table C-14 (continued).

Site	Price Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
AP-36	0.0001	NA	NA
AP-37	0.0005	NA	NA
AP-38	0.0093	NA	NA
AP-39	0.0003	NA	NA
AP-40	0.0003	0.0007	NA
AP-41	7.0E-05	NA	NA
AP-42	0.0010	NA	NA
AP-43	0.0026	NA	NA
AP-45	0.0002	NA	NA
AP-46	0.0006	NA	NA
AP-47	6.8E-05	NA	NA
GC-1	0.0010	0.0025	0.0038
GC-2	0.0002	0.0006	0.0010
GC-3	0.0122	NA	NA
GC-4	0.0002	0.0006	0.0009
GC-5	0.0011	0.0027	0.0041
GC-6	0.0007	0.0017	0.0026
GC-7	0.0015	NA	NA
GC-8	0.0009	NA	NA
GC-9	2.3E-05	5.7E-05	8.7E-05
GC-10	0.0006	NA	NA
GC-11	0.0003	NA	NA
GC-12	0.0007	NA	NA
GC-13	0.0018	NA	NA
GC-14	0.0004	0.0009	0.0014
GC-15	0.0008	NA	NA
GC-16	0.0005	0.0012	0.0019
GC-17	0.0011	NA	NA
GC-18	0.0003	NA	NA
GC-19	0.0012	NA	NA
GC-20	0.0011	NA	NA
GC-21	0.0002	NA	NA
GC-22	0.0053	NA	NA
GC-23	0.0001	NA	NA
GC-24	0.0005	0.0012	0.0018
GC-25	0.0293	0.0721	NA

Table C-14 (continued).

Site	Price Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
GC-26	0.0004	0.0009	0.0013
GC-27	0.0069	0.0168	NA
GC-28	0.0003	NA	NA
GC-29	0.0039	NA	NA
GC-30	0.0015	NA	NA
GC-31	8.5E-05	0.0002	0.0003
GC-32	0.0008	NA	NA
GC-33	0.0013	NA	NA
GC-34	0.0007	NA	NA
GC-35	0.0050	NA	NA
GC-36	0.0266	NA	NA
GC-37	3.7E-05	9.2E-05	0.0001
GC-38	0.0010	NA	NA
GC-39	0.0151	NA	NA
GC-40	0.0111	NA	NA
GC-41	0.1082	NA	NA
GC-42	0.0121	NA	NA
GC-43	0.0059	NA	NA
GC-44	0.0005	0.0012	NA
GC-45	0.0160	0.0397	NA
GC-46	0.0012	0.0030	0.0047
GC-47	0.0177	0.0435	NA
GC-48	0.0060	NA	NA
GC-49	0.0196	0.0482	NA
GC-50	0.0004	NA	NA
GC-51	0.0163	0.0397	NA
GC-52	0.0007	NA	NA
GC-53	0.0071	NA	NA
GC-54	5.3E-05	0.0001	0.0002
GC-55	0.0003	NA	NA
GC-56	0.0012	NA	NA
GC-57	0.0003	NA	NA
GC-58	0.0049	NA	NA
MC-1	0.0037	0.0093	0.0136
MC-4	0.0015	0.0040	NA
MC-5	0.2312	0.5809	NA

Table C-14 (continued).

Site	Price Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
MC-6	0.0089	0.0225	0.0330
MC-7	0.0062	NA	NA
MC-8	0.0056	NA	NA
MC-9	0.0216	NA	NA
MC-10	0.0006	NA	NA
MC-11	0.0072	NA	NA
MC-12	0.0046	0.0114	0.0168
MC-13	0.0216	NA	NA
MC-14	0.0136	0.0342	NA
MC-16	0.0040	0.0100	0.0147
MC-17	0.0056	NA	NA
MC-18	0.0022	NA	NA
MC-19	0.0062	NA	NA
MC-20	0.0019	0.0048	0.0071
MC-21	0.0014	0.0034	0.0051
MC-22	0.0097	NA	NA
MC-23	0.0008	NA	NA
MC-24	0.0045	NA	NA
MC-25	0.0012	NA	NA
MC-26	0.0034	NA	NA
RM-2	0.0003	0.0008	0.0013
RM-3	0.0018	0.0045	0.0070
RM-4	8.1E-05	0.0002	0.0003
RM-5	0.0213	0.0532	0.0826
RM-6	0.0280	0.0698	0.1084
RM-7	0.0260	0.0651	0.1006
RM-8	0.0004	0.0010	0.0015
RM-9	0.0002	0.0005	0.0007
RM-10	0.0020	0.0051	0.0079
RM-11	4.1E-05	0.0001	0.0002
RM-12	0.0003	0.0007	0.0011
RM-13	3.9E-06	9.8E-06	1.5E-05
RM-14	0.0064	0.0160	0.0248
RM-15	0	0	0
RM-16	0.0011	0.0027	0.0041
RM-17	0.0008	0.0020	0.0030

Table C-14 (continued).

Site	Price Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
RM-18	0.0007	0.0017	0.0026
RM-19	0.0008	0.0021	0.0032

C.7 Sensitivity Analysis: Ratio of methane (routine) emissions allocated to each product (Natural Gas, Natural Gas Liquids (NGL), and Oil) divided by production for an assumed well lifetime of 15 years

Tables C-15 to C-17 shows the ratio of total emissions to produced components when a well lifetime of 15 years is assumed instead of the 30 year lifetime.

Table C-15. Ratio of methane (routine) emissions allocated to each salable product (Natural Gas, Natural Gas Liquids, and Oil) divided by its respective estimated ultimate recovery (15 years well lifetime), based on a **mass allocation**. The values represent the ratio of the sum of emissions divided by the sum of production across all sites where each specific component is produced.

	Pneumatic Controllers	Pneumatic Pumps	Equipment Leaks	Total
Emitted Methane (scf) / Produced Natural Gas (scf)	0.0005	0.0001	0.0001	0.0007
Emitted Methane (Mg) / Produced Natural Gas Liquids (Mg)	0.0009	1.4E-05	9.2E-05	0.0010
Emitted Methane (Mg) / Produced Oil (Mg)	0.0007	1.2E-06	6.7E-05	0.0008

Table C-16. Ratio of methane (routine) emissions allocated to each salable product (Natural Gas, Natural Gas Liquids, and Oil) divided by its respective estimated ultimate recovery (15 years well lifetime), based on an **energy allocation**. The values represent the ratio of the sum of emissions divided by the sum of production across all sites where each specific component is produced.

	Pneumatic Controllers	Pneumatic Pumps	Equipment Leaks	Total
Emitted Methane (scf) / Produced Natural Gas (scf)	0.0005	0.0001	0.0001	0.0007
Emitted Methane (Mg) / Produced Natural Gas Liquids (Mg)	0.0009	1.3E-05	8.9E-05	0.0010
Emitted Methane (Mg) / Produced Oil (Mg)	0.0007	1.1E-06	6.4E-05	0.0007

Table C-17. Ratio of methane (routine) emissions allocated to each salable product (Natural Gas, Natural Gas Liquids, and Oil) divided by its respective estimated ultimate recovery (15 years well lifetime), based on a **price allocation**. The values represent the ratio of the sum of emissions divided by the sum of production across all sites where each specific component is produced.

	Pneumatic Controllers	Pneumatic Pumps	Equipment Leaks	Total
Emitted Methane (scf) / Produced Natural Gas (scf)	0.0005	0.0001	0.0001	0.0007
Emitted Methane (Mg) / Produced Natural Gas Liquids (Mg)	0.0011	2.9E-05	0.0001	0.0012
Emitted Methane (Mg) / Produced Oil (Mg)	0.0010	2.5E-06	9.8E-05	0.0011

C.8 *Sensitivity Analysis: Ratio of methane emissions from routine production allocated to each product (Natural Gas, Natural Gas Liquids (NGL), and Oil) divided by production when daily production at the time of measurement is used instead of estimated ultimate recovery.*

Tables C-18 to C-20 show the ratio of total emissions to produced components, when daily production at the time of the measurement is used instead of estimated ultimate recovery.

Table C-18. Ratio of methane daily emissions allocated to each salable product (Natural Gas, Natural Gas Liquids, and Oil) divided by its respective daily production, based on a **mass allocation**. The values represent the ratio of the sum of emissions divided by the sum of production across all sites where each specific product is produced.

	Pneumatic Controllers	Pneumatic Pumps	Equipment Leaks	Total
Emitted Methane (scf) / Produced Natural Gas (scf)	0.0002	5.1E-05	2.0E-06	0.0003
Emitted Methane (Mg) / Produced Natural Gas Liquids (Mg)	0.0002	5.7E-06	8.4E-07	0.0002
Emitted Methane (Mg) / Produced Oil (Mg)	7.3E-05	2.4E-07	2.9E-07	7.4E-05

Table C-19. Ratio of methane daily emissions allocated to each salable product (Natural Gas, Natural Gas Liquids, and Oil) divided by its respective daily production, based on an **energy allocation**. The values represent the ratio of the sum of emissions divided by the sum of production across all sites where each specific product is produced.

	Pneumatic Controllers	Pneumatic Pumps	Equipment Leaks	Total
Emitted Methane (scf) / Produced Natural Gas (scf)	0.0002	5.1E-05	2.0E-06	0.0003
Emitted Methane (Mg) / Produced Natural Gas Liquids (Mg)	0.0002	5.2E-06	8.2E-07	0.0002
Emitted Methane (Mg) / Produced Oil (Mg)	7.2E-05	2.3E-07	2.9E-07	7.2E-05

Table C-20. Ratio of methane daily emissions allocated to each salable product (Natural Gas, Natural Gas Liquids, and Oil) divided by its respective daily production, based on a **price allocation**. The values represent the ratio of the sum of emissions divided by the sum of production across all sites where each specific product is produced.

	Pneumatic Controllers	Pneumatic Pumps	Equipment Leaks	Total
Emitted Methane (scf) / Produced Natural Gas (scf)	0.0002	5.1E-05	1.9E-06	0.0002
Emitted Methane (Mg) / Produced Natural Gas Liquids (Mg)	0.0002	1.2E-05	9.7E-07	0.0002
Emitted Methane (Mg) / Produced Oil (Mg)	8.5E-05	3.1E-07	3.4E-07	8.8E-05

C.9 Allocation of Emissions from Completion Flowbacks

Table C-21. Measured methane emissions and total production for each completion event where measurements were made by Allen, *et al.*, [8] and the host company reported production of gas and oil.

Event	Methane Emissions (Mg)	Gas EUR (Mg) (30-yr)	Oil EUR (Mg) (30-yr)	Gas EUR (Mg) (15-yr)	Oil EUR (Mg) (15-yr)
AP1	0.4992	228.5	0	177.8	0
AP2	0.1229	137,079	0	106,695	0
AP3	1.824	155,356	0	120,921	0
GC1	2.016	93,150	166,883	72,503	129,894
GC2	1.728	83,835	159,245	65,253	123,948
GC3	5.376	40,365	13,187	31,418	10,264
GC4	3.456	24,840	11,321	19,334	8,812
GC5	0.3322	117,441	0	91,410	0
GC6	4.742	189,404	1,851	147,423	1,441
GC7	1.728	46,575	116,167	36,252	90,419
MC1	3.840	30,592	10,465	23,812	8,145
MC2	0.5184	30,592	20,001	23,812	15,568
MC3	0.0518	70,334	0	54,744	0
MC4	0.0461	85,719	0	66,720	0
MC5	0.0403	83,521	0	65,009	0
RM3	0.1997	8,884	42,210	6,915	32,854
RM4	0.5760	8,884	47,599	6,915	37,049
RM5	0.7488	11,845	51,529	9,220	40,108
RM6	0.6528	2,961	3,983	2,305	3,100

Table C-22. Completion flowback emissions. Based on a **mass allocation**, fraction of production for each salable product and methane emissions allocated to each product.

Event	Natural Gas, fraction of total production (mass basis)	Natural Gas Liquids, fraction of total production (mass basis)	Oil, fraction of total production (mass basis)	Emissions allocated to Natural Gas (Mg)	Emissions allocated to Natural Gas Liquids (Mg)	Emissions allocated to Oil (Mg)
AP1	1	0	0	0.4992	0	0
AP2	1	0	0	0.1229	0	0
AP3	1	0	0	1.8240	0	0
GC1	0.20	0.16	0.64	0.4061	0.3161	1.2938
GC2	0.19	0.15	0.66	0.3351	0.2609	1.1320
GC3	0.42	0.33	0.25	2.2785	1.7737	1.3238
GC4	0.39	0.30	0.31	1.3349	1.0391	1.0820
GC5	1	0	0	0.3322	0	0
GC6	0.56	0.43	0.01	2.6408	2.0557	0.0459
GC7	0.16	0.13	0.71	0.2781	0.2165	1.2335
MC1	0.68	0.07	0.25	2.6006	0.2607	0.9787
MC2	0.55	0.06	0.40	0.2849	0.0286	0.2049
MC3	1	0	0	0.0518	0	0
MC4	1	0	0	0.0461	0	0
MC5	1	0	0	0.0403	0	0
RM3	0.10	0.07	0.83	0.0209	0.0138	0.1650
RM4	0.09	0.06	0.84	0.0545	0.0361	0.4854
RM5	0.11	0.07	0.81	0.0842	0.0558	0.6088
RM6	0.26	0.17	0.57	0.1674	0.1110	0.3744

Table C-23. Completion flowback emissions. Based on an **energy allocation**, fraction of production for each salable product and methane emissions allocated to each product.

Event	Natural Gas, fraction of total production (energy basis)	Natural Gas Liquids, fraction of total production (energy basis)	Oil, fraction of total production (energy basis)	Emissions allocated to Natural Gas (Mg)	Emissions allocated to Natural Gas Liquids (Mg)	Emissions allocated to Oil (Mg)
AP1	1	0	0	0.4992	0	0
AP2	1	0	0	0.1229	0	0
AP3	1	0	0	1.8240	0	0
GC1	0.22	0.16	0.62	0.4441	0.3153	1.2566
GC2	0.21	0.15	0.64	0.3669	0.2605	1.1007
GC3	0.45	0.32	0.23	2.4151	1.7147	1.2462
GC4	0.41	0.29	0.30	1.4223	1.0098	1.0239
GC5	1	0	0	0.3322	0	0
GC6	0.58	0.41	0.01	2.7486	1.9514	0.0424
GC7	0.18	0.13	0.70	0.3059	0.2172	1.2050
MC1	0.71	0.06	0.23	2.7177	0.2476	0.8747
MC2	0.59	0.05	0.36	0.3038	0.0277	0.1869
MC3	1	0	0	0.0518	0	0
MC4	1	0	0	0.0461	0	0
MC5	1	0	0	0.0403	0	0
RM3	0.12	0.07	0.82	0.0230	0.0139	0.1629
RM4	0.10	0.06	0.83	0.0600	0.0362	0.4798
RM5	0.12	0.07	0.80	0.0925	0.0558	0.6005
RM6	0.28	0.17	0.56	0.1808	0.1091	0.3629

Table C-24. Completion flowback emissions. Based on a **price allocation**, fraction of production for each salable product and methane emissions allocated to each product.

Event	Natural Gas, fraction of total production (price basis)	Natural Gas Liquids, fraction of total production (price basis)	Oil, fraction of total production (price basis)	Emissions allocated to Natural Gas (Mg)	Emissions allocated to Natural Gas Liquids (Mg)	Emissions allocated to Oil (Mg)
AP1	1	0	0	0.4992	0	0
AP2	1	0	0	0.1229	0	0
AP3	1	0	0	1.8240	0	0
GC1	0.06	0.13	0.81	0.1298	0.2603	1.6259
GC2	0.06	0.12	0.82	0.1061	0.2127	1.4092
GC3	0.19	0.38	0.43	1.0164	2.0380	2.3217
GC4	0.16	0.32	0.51	0.5582	1.1192	1.7787
GC5	1	0	0	0.3322	0	0
GC6	0.33	0.65	0.02	1.5430	3.0939	0.1054
GC7	0.05	0.10	0.85	0.0845	0.1695	1.4740
MC1	0.37	0.10	0.53	1.4318	0.3684	2.0398
MC2	0.25	0.06	0.68	0.1302	0.0335	0.3546
MC3	1	0	0	0.0518	0	0
MC4	1	0	0	0.0461	0	0
MC5	1	0	0	0.0403	0	0
RM3	0.03	0.05	0.92	0.0059	0.0100	0.1838
RM4	0.03	0.04	0.93	0.0151	0.0258	0.5351
RM5	0.03	0.05	0.91	0.0238	0.0406	0.6844
RM6	0.09	0.15	0.77	0.0563	0.0960	0.5005

Table C-25. Ratio of methane emissions from completion flowback events allocated to each salable product (Natural Gas, Natural Gas Liquids, and Oil) divided by its estimated ultimate production (EUR), assuming **30 year well-lifetime**, based in a **mass allocation**.

Mass Basis			
Event	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
AP1	0.0022	NA	NA
AP2	9.2E-07	NA	NA
AP3	1.2E-05	NA	NA
GC1	8.0E-06	7.8E-06	7.8E-06
GC2	7.3E-06	7.1E-06	7.1E-06
GC3	0.0001	0.0001	0.0001
GC4	9.8E-05	9.6E-05	9.6E-05
GC5	2.9E-06	NA	NA
GC6	2.5E-05	2.5E-05	2.5E-05
GC7	1.1E-05	1.1E-05	1.1E-05
MC1	9.6E-05	9.4E-05	9.4E-05
MC2	1.1E-05	1.0E-05	1.0E-05
MC3	7.5E-07	NA	NA
MC4	5.4E-07	NA	NA
MC5	4.9E-07	NA	NA
RM3	4.0E-06	3.9E-06	3.9E-06
RM4	1.0E-05	1.0E-05	1.0E-05
RM5	1.2E-05	1.2E-05	1.2E-05
RM6	9.7E-05	9.4E-05	9.4E-05

Table C-26. Ratio of methane emissions from completion flowback events allocated to each salable product (Natural Gas, Natural Gas Liquids, and Oil) divided by its estimated ultimate production (EUR), assuming **30 year well-lifetime**, based in an **energy allocation**

Event	Energy Basis		
	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
AP1	0.0022	NA	NA
AP2	9.2E-07	NA	NA
AP3	1.2E-05	NA	NA
GC1	8.7E-06	7.7E-06	7.5E-06
GC2	8.0E-06	7.1E-06	6.9E-06
GC3	0.0001	9.7E-05	9.5E-05
GC4	0.0001	9.3E-05	9.0E-05
GC5	2.9E-06	NA	NA
GC6	2.6E-05	2.4E-05	2.3E-05
GC7	1.2E-05	1.1E-05	1.0E-05
MC1	0.0001	8.9E-05	8.4E-05
MC2	1.1E-05	9.9E-06	9.3E-06
MC3	7.5E-07	NA	NA
MC4	5.4E-07	NA	NA
MC5	4.9E-07	NA	NA
RM3	4.4E-06	3.9E-06	3.9E-06
RM4	1.2E-05	1.0E-05	1.0E-05
RM5	1.3E-05	1.2E-05	1.2E-05
RM6	0.0001	9.2E-05	9.1E-05

Table C-27. Ratio of methane emissions from completion flowback events allocated to each salable product (Natural Gas, Natural Gas Liquids, and Oil) divided by its estimated ultimate production (EUR), assuming **30 year well-lifetime**, based in a **price allocation**.

Price Basis			
Event	Emitted Methane (scf) / EUR Natural Gas (scf)	Emitted Methane (Mg) / EUR Natural Gas Liquids (Mg)	Emitted Methane (Mg) / EUR Oil (Mg)
AP1	0.0022	NA	NA
AP2	9.2E-07	NA	NA
AP3	1.2E-05	NA	NA
GC1	2.5E-06	6.4E-06	9.7E-06
GC2	2.3E-06	5.8E-06	8.8E-06
GC3	4.6E-05	0.0001	0.0002
GC4	4.1E-05	0.0001	0.0002
GC5	2.9E-06	NA	NA
GC6	1.5E-05	3.7E-05	5.7E-05
GC7	3.3E-06	8.3E-06	1.3E-05
MC1	5.3E-05	0.0001	0.0002
MC2	4.8E-06	1.2E-05	1.8E-05
MC3	7.5E-07	NA	NA
MC4	5.4E-07	NA	NA
MC5	4.9E-07	NA	NA
RM3	1.1E-06	2.8E-06	4.4E-06
RM4	2.9E-06	7.3E-06	1.1E-05
RM5	3.4E-06	8.6E-06	1.3E-05
RM6	3.2E-05	8.1E-05	0.0001

C.10 Rationale for disaggregation of emission sources not directly measured.

Table C-28. Additional sources of methane emissions from natural gas production considered in the EPA GHG national inventory [9]. Rationale for disaggregation of emissions into NG = Natural Gas, NGL= Natural Gas Liquids, and Oil is provided. The column showing net emissions refers to total emissions from each category before the allocation to the corresponding products.

EPA GHG Inventory Activity	Net Emissions (Gg methane/yr)	Classification	Rationale for classification
Refractures	143	(NG+NGL+OIL)	Emission from well associated with all products from well.
Gas wells without HF	13	(NG+NGL+OIL)	Emission from well associated with all products from well
Gas wells with HF	15	(NG+NGL+OIL)	Emission from well associated with all products from well
Separators	57	(NG+NGL+OIL)	Emission from well associated with all products from well
Meters/Piping	54	(NG+NGL+OIL)	In the absence of specific information, assume these devices handle all products.
Heaters	18	(NG+NGL+OIL)	In the absence of specific information, assume these devices handle all products.
Dehydrators	16	(NG+NGL)	Device that handles exclusively gas from the well site, which are separated into NG and NGL products.
Workovers without HF	0.3	(NG+NGL+OIL)	Emission from well associated with all products from well
Liquids Unloading (without plunger lifts)	149	(NG+NGL+OIL)	Emission from well associated with all products from well
Liquids Unloading (with plunger lifts)	108	(NG+NGL+OIL)	Emission from well associated with all products from well
Kimray Pumps	185	(NG+NGL)	Device that handles exclusively gas.
Condensate Tanks without Controls	94	(OIL only)	Condensate tanks handle OIL product.

Table C-28 (continued).

EPA GHG Inventory Activity		Net Emissions (Gg methane/yr)	Classification	Rationale for classification
Condensate with Controls	Tanks	52	(OIL only)	Condensate tanks handle OIL product.
Gas Engines		227	(NG+NGL)	Device that handles exclusively gas.
Dehydrators Vents		41	(NG+NGL)	Device that handles exclusively gas.
Small Reciprocating Compressors			(NG+NGL)	Device that handles exclusively gas.
Large Reciprocating Compressors		49	(NG+NGL)	Device that handles exclusively gas.
Large Reciprocating Stations			(NG+NGL)	Device that handles exclusively gas.
Pipeline Leaks		90	(NG+NGL)	Pipelines handle gas products.
Well Drilling		0.4	(NG+NGL+OIL)	Emission from well associated with all products from well
Vessel Blowdowns		0.4	(NG+NGL)	Involves venting of gas products.
Pipeline Blowdowns		2	(NG+NGL)	Involves venting of gas products.
Compressor Blowdowns		2	(NG+NGL)	Involves venting of gas products.
Compressor Starts		3	(NG+NGL)	Involves venting of gas products.
Pressure Relief Valves		0.4	(NG+NGL)	Involves venting of gas products.
Mishaps		1	(NG+NGL+OIL)	In the absence of specific information, assume these events are associated with all products.

C.11 References

- (1) API. (2009). Compendium of Greenhouse Gas Emissions Methodologies for Oil and Natural Gas Industry. Washington, DC: American Petroleum Institute.
- (2) American Gas Association. (1951). Fuel Flue Gases.
- (3) GREET, developed by Argonne National Laboratory, from <http://greet.es.anl.gov/>
- (4) EIA. (2012). Annual Energy Outlook. DOE/EIA.
- (5) EIA. (2013). Oil and Gas Supply Module of the National Energy Modeling System: Model Documentation 2013, Appendix 2.c. Retrieved 1, 14, 2014, from [http://www.eia.gov/forecasts/aeo/nems/documentation/ogsm/pdf/m063\(2013\).pdf](http://www.eia.gov/forecasts/aeo/nems/documentation/ogsm/pdf/m063(2013).pdf)
- (6) O'Sullivan, F., & Paltsev, S. (2012). Shale gas production: potential versus actual greenhouse gas emissions. *Environmental Research Letters*, 7, 1-7.
- (7) NETL. 2012. Role of Alternative Energy Sources: Natural Gas Technology Assessment. U.S. Department of Energy.
- (8) Allen DT, Torres VM, Thomas J, Sullivan D, Harrison M, et al. 2013. Measurements of Methane Emissions at Natural Gas Production Sites: Study Appendices and Database, Austin, Texas: University of Texas <http://dept.ceer.utexas.edu/methane/study/>
- (9) EPA. (2013, April 12). Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2011. Retrieved 4 22, 2013, from <http://www.epa.gov/climatechange/ghgemissions/usinventoryreport.html>

D Criteria and Air Toxics Emissions from Shale Gas Production

D.1 Hydrocarbon concentrations at Flower Mound Shiloh and Hinton monitoring sites

The concentrations of light alkanes at EML are higher than at the other two sites. At EML, the average morning maximum concentrations of ethane, propane, and butane total close to 100 ppbC, while at Flower Mound Shiloh and Hinton, the morning totals are 60 ppbC and 40 ppbC respectively. Summed over all days and hours in the 20 month sampling period, the average concentrations at EML for ethane, propane, and butane are 31.3, 19.4, and 9.2 ppbC, respectively. For Flower Mound Shiloh those averages are 18.4, 13.0, and 7.8, respectively. In the case of Hinton, the averages are 15.5 for ethane, 11.0 for propane, and 6.5 for butane.

Figures D-1 and D-2 show daily average hydrocarbon concentrations at Flower Mound Shiloh and Hinton, respectively. The dominant species (ethane, propane and butane) are similar to EML, since, in addition to being associated with emissions from natural gas production, these species are commonly found in regional background hydrocarbon samples.

Figure D-3 shows average diurnal hydrocarbon concentrations at Flower Mound Shiloh for observations with winds between 135° and 225°, the wind directions for which Flower Mound Shiloh was used as a background site.

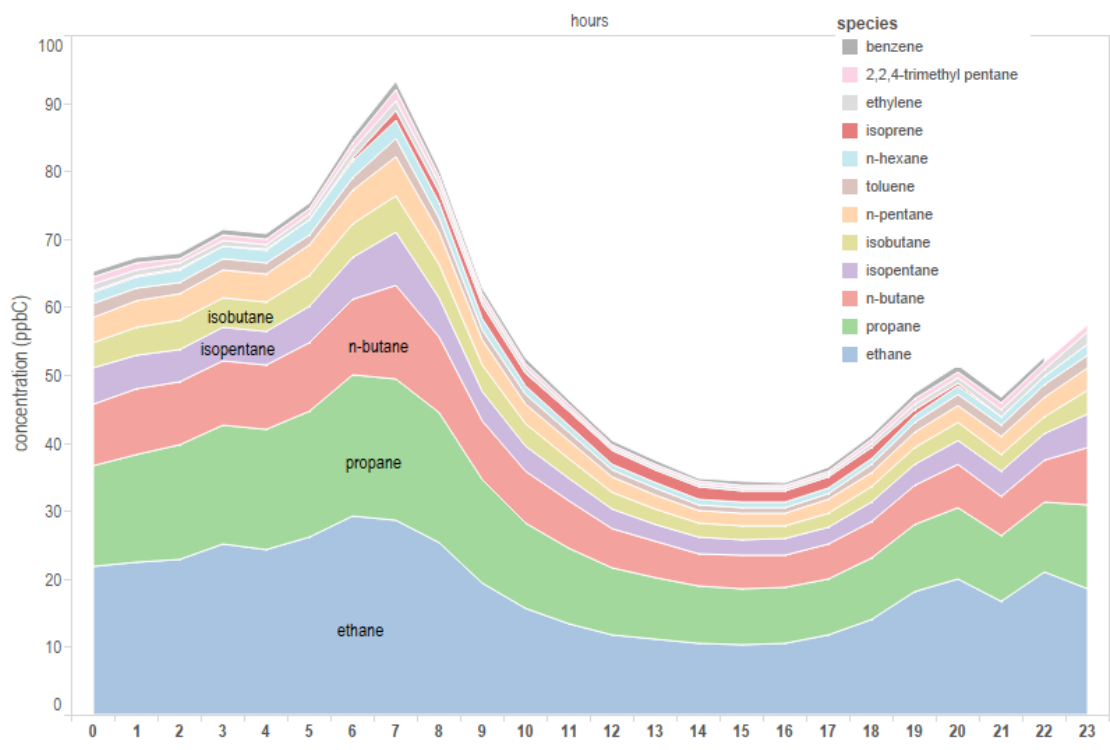


Figure D-1. Diurnal pattern of hydrocarbon concentrations (ppbC) at Flower Mound Shiloh averaged over 20 months of sampling.

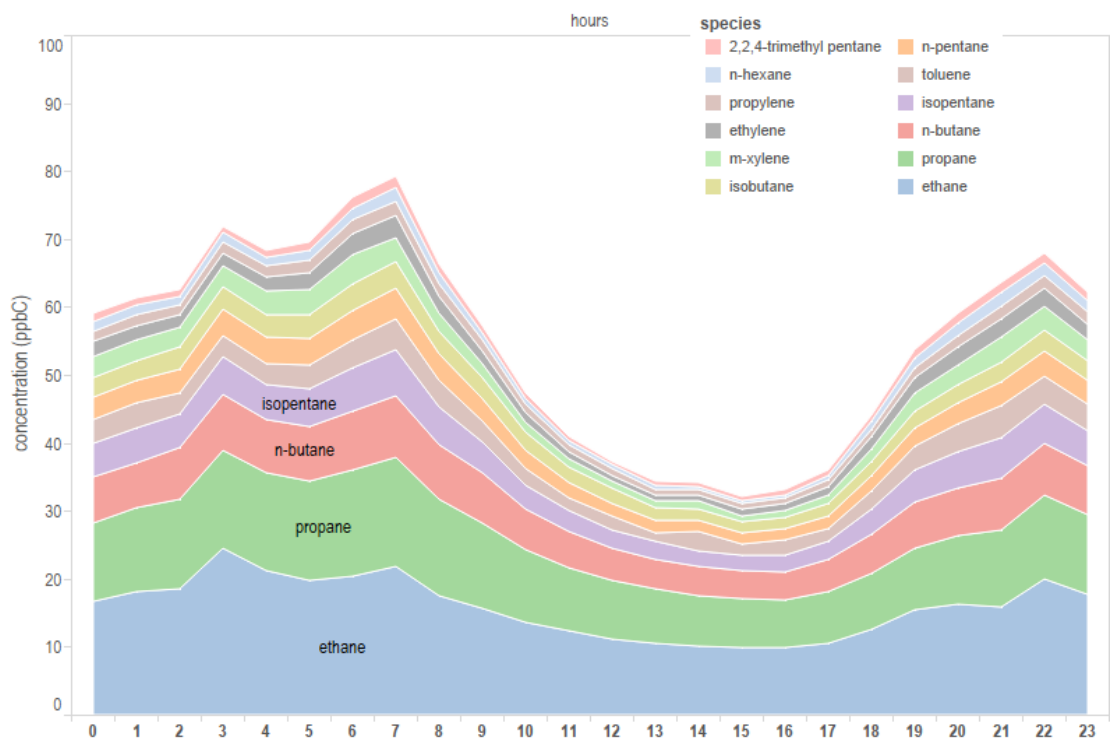


Figure D-2. Diurnal pattern of hydrocarbon concentrations (ppbC) at Hinton averaged over 20 months of sampling.

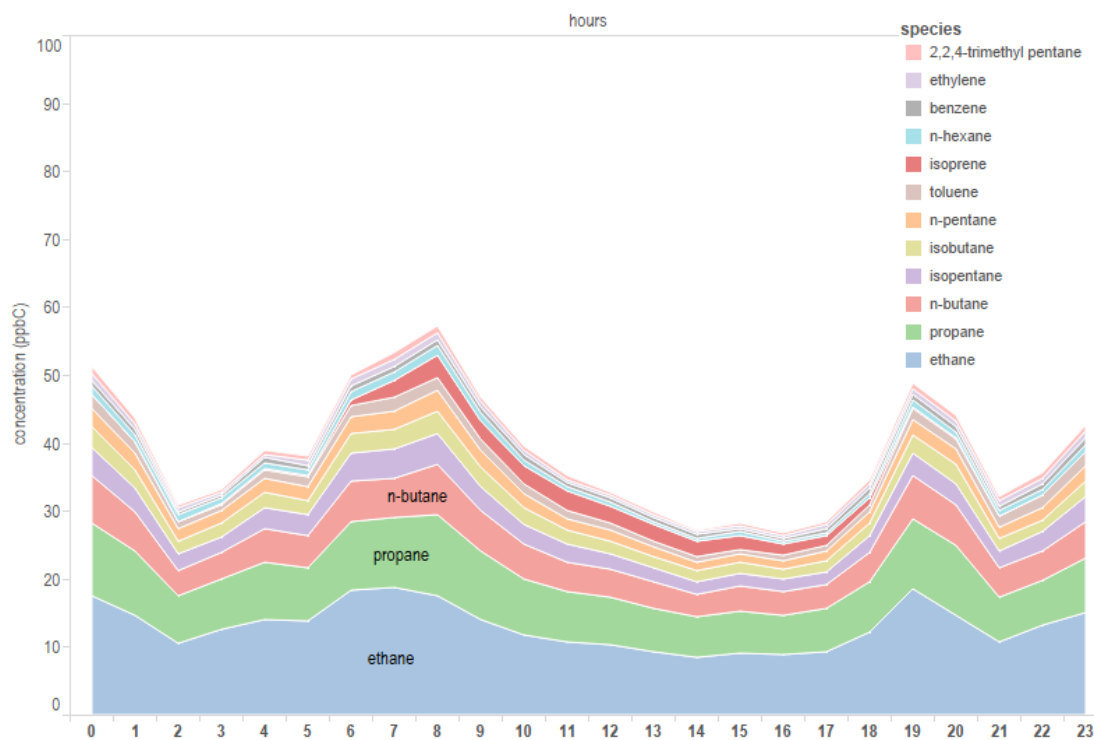


Figure D-3. Average diurnal pattern of hydrocarbon concentrations at Flower Mound Shiloh filtering for hours when wind direction was between 135 ° – 225 °.

D.2 Comparison of measurements and model predictions when Hinton is used as background site but data is filtered for the same hours considered in the Flower Mound Shiloh dataset

Figure D-4 shows a comparison between predicted VOC concentrations calculated using the AERMOD dispersion model, and the observed, background corrected alkane concentrations as they were measured at the EML monitoring site. When Hinton was used as a background site, all directions were considered, however, as a sensitivity test, the present analysis reflects only those hours considered in the final dataset of Flower Mound Shiloh (filtering for wind directions with the least influence from natural gas sources for that particular monitoring site).

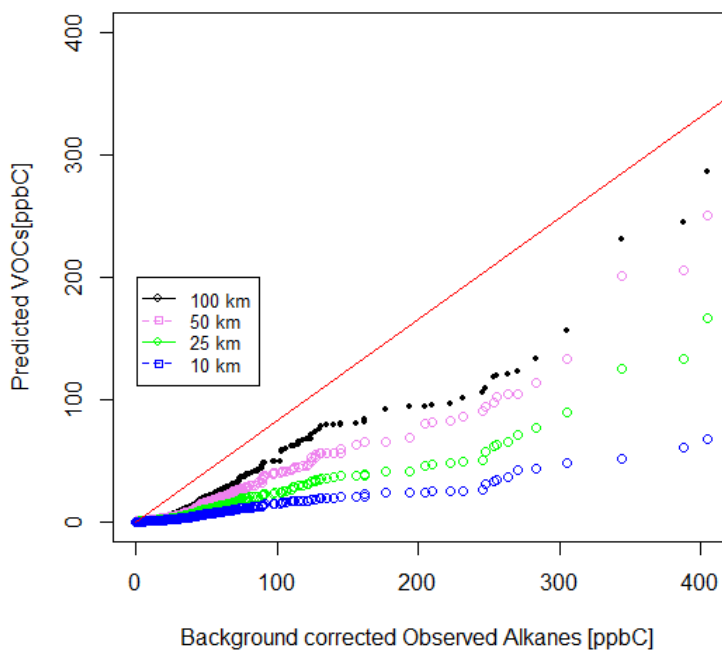


Figure D-4. Predicted total VOC concentrations vs. background corrected observed summed alkane concentrations at EML, when *Hinton* was used as background site, filtering for the same hours that were considered in the Flower Mound Shiloh final dataset. For four sets of dispersion calculations, within 10, 25, 50, and 100 km radii of the observation site. The red line has a slope of 0.83 since 83% of the total VOC are speciated as alkanes.

The regression has a slope of 0.55 (standard error 0.009), for observations over 50 ppbC, as compared to a slope of 0.66 when Flower Mound Shiloh is used as background site.

D.3 Calculation of emissions from pneumatic devices, chemical injection pumps, and equipment leaks based on Allen, et al (2013)

Table D-1 shows the emission factors used to estimate emissions from pneumatic devices, chemical injection pumps, and equipment leaks from production sites. For estimated emissions of propane and higher alkanes, a weighted average gas composition for the Barnett Shale was used (89.16% methane, 4.24% ethane; 2.67% C₃+ alkanes

(molar basis; weighted by total gas production for each site), or 77.37% methane, and 7.45% C₃₊ alkanes (mass basis)).

Table D-1. Routine emissions from production sites.

	Methane Scf/min per device*	Activity factor for the Barnett Shale**	Methane Scf/min per well	Methane Mg/yr per well***	C₃₊ Mg/yr per well †
Pneumatic Devices	0.175	8,086 devices	0.507	5.116	0.493
Chemical Injection Pumps	0.192	1,972 devices	0.136	1.369	0.132
Equipment Leaks (per well)	0.064	2,791 wells	0.064	0.646	0.062
Total	0.431		0.707	7.131	0.687

* Emission factors were obtained from the Supporting information of Allen, *et al* (2013), Tables S2-2, 3, 4.

** Activity factors for the Barnett Shale were calculated from total counts (for the host companies that participated in the study) for the play (based on AAPG basin classification) from the database from Allen *et al*'s.

*** 1scf of methane = 19.2 g of methane.

† Mg/yr per well of propane and higher alkanes are obtained by multiplying Methane Mg/yr per well by the mass ratio of C₃₊ to methane (7.45%/77.37% = 0.096).

The total methane emissions of 7.13 Mg per year per well from pneumatic devices, chemical injection pumps, and equipment leaks, result in 0.687 Mg/yr per well of propane and higher alkanes, multiplied by the total number of wells reported by the TCEQ for the year 2011; 14,886 the total emissions would be roughly 10,200 tons per year of propane and higher alkanes.

E Developing a Methane, Ethane, and Propane Emission Inventory for the Barnett Shale Production Region

E.1 Spatial interpolation of percent VOC

For the 8,025 sites with reported VOC emissions in the TCEQ special inventory, only 30% reported %VOC values. Figure E-1 shows the spatial distribution of sites with known %VOC and with unknown %VOC. Assuming that %VOC and gas composition are well characterized by the spatial location of sites, it is expected that the missing values can be estimated from neighboring sites with known gas composition using an inverse distance weight (IDW) method for interpolation. Figure E-2 shows the spatially resolved distribution of grid cells with interpolated %VOC values.

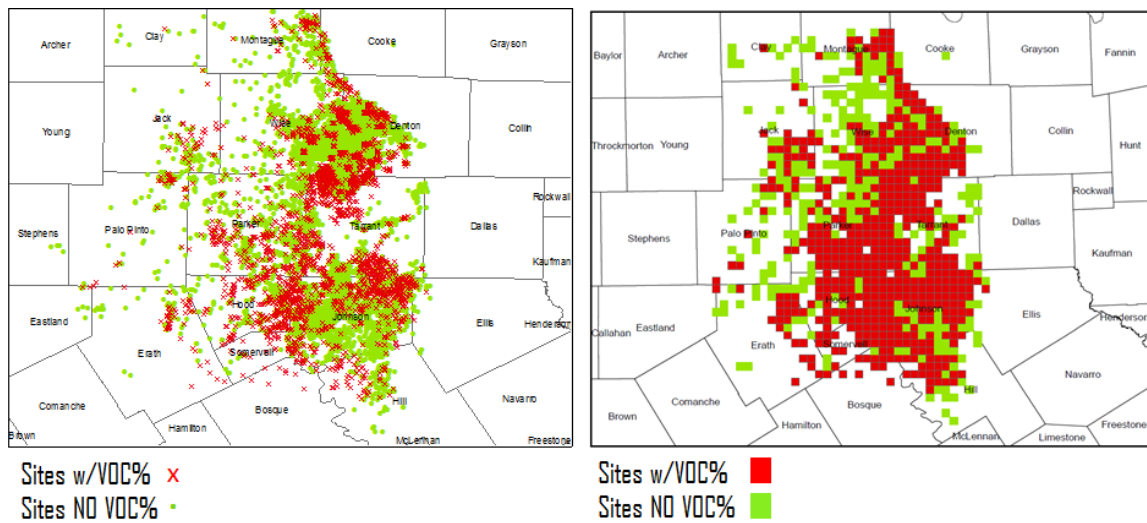


Figure E-1. Location of sites from the TCEQ special inventory, with known, and unknown %VOC.

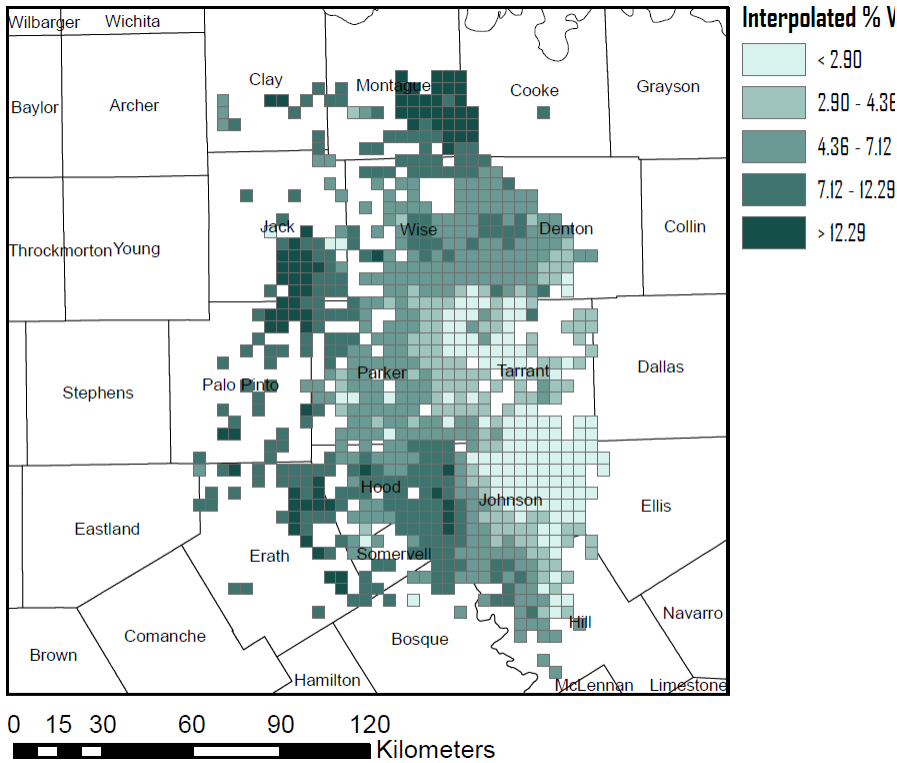


Figure E-2. Spatially resolved distribution of interpolated % VOC for grid cells that contain Natural Gas production sites. Coloring scheme shows the values of % VOC for each grid cell.

E.2 Detailed methodology for the estimation of emissions by source category

Condensate and oil Tanks: Correlations developed by Vazquez and Beggs [1] were used to estimate dissolved gas in oil. The correlations estimate gas solubility based on temperature, pressure, and API gravity. Average conditions for separators feeding liquids to tanks were determined (70 psig, 40° API) [2] from data collected at gas wells in the Barnett Shale, yielding an estimated dissolved gas loading of 80 scf/bbl of condensate [2, 3]. The dissolved gas loading is multiplied by the site-specific condensate and oil production to yield the volume of dissolved gas vented per unit of time. To estimate methane, ethane, and propane emissions, it is assumed that these three species are the only species that flash, and that the relative composition of the three species in the vented gas is similar to the produced gas. Site-specific estimated %C1, %C2, %C3 values are used to determine vented emissions.

Water Tanks: For Water Tanks the solubility of methane, ethane, and propane in water was estimated based on Henry's law [2, 4] and site specific produced water production rate.

Because the solubility of methane, ethane, and propane in water depends on separator conditions, average temperature and pressure for a separator were determined (70 psig, 60° F) from data collected at production sites in the Barnett Shale [2]. Combining the average temperature and pressure with site-specific gas composition, partial pressures of methane, ethane and propane above the water tank were computed using Henry's law. Partial pressures were multiplied by the amount of water per barrel (8830 moles/bbl), density of the species, and site-specific water production rates to yield C1, C2, and C3 emissions.

Loadings: It was assumed that loadings of liquids from tanks to vehicles would generate no additional methane, ethane, propane emissions besides those that were already accounted for in the tanks source category.

Frac Tanks: Allen, *et al.* [2] reported methane emission rates for 5 completion flowback events in the Mid Continent region (where the Barnett Shale production region is situated), with an average regional emission rate of 2.8 MMscf/yr . It is assumed that methane emissions from frac tanks are equal to those emissions, and ethane, propane emissions are estimated based on the site-specific gas composition.

All other source categories: For the rest of the source categories, it is assumed that methane, ethane, and propane emissions can be scaled directly from the VOC emissions, based on the gas composition. A special consideration was taken for fugitive emissions. In a previously published work [5] a correction factor was developed in a comparison between the TCEQ special inventory and ambient VOC concentrations. Fugitive emissions are corrected for their underestimation in the special inventory based on emission rates estimated from direct measurements [2].

VOC emissions from pneumatic devices, chemical injection pumps, and equipment leaks, are 0.021 scf/min [2], or 0.687 Mg/yr per well. For a count of 14,886 producing wells in the Barnett Shale [5], VOC emissions from those sources would be roughly 10,200 tons per year. This is 159% higher than the corresponding estimate for similar categories in the TCEQ inventory (3,935 tons per year for fugitives).

Emissions were estimated for each source category, then summed by site to spatially locate them, and finally summed up by grid cell. To determine the corresponding methane, ethane, propane emissions from VOC emissions without source category, VOC emissions were spatially distributed among the rest of the sites (weighted by the magnitude of VOC emissions) and methane, ethane, propane emissions were also scaled based on VOC emissions and the gas composition of the site where they were assigned.

E.3 Estimated methane emissions by grid cell

Table E-1 shows final data set of grid cells, with location and production data.

Table E-1. Location and production data for each grid cell in the final data set.

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
1066	31	15	0.1	0.0
1139	30	16	313.8	300.8
1285	28	18	106.1	878.5
1287	30	18	149.6	229.2
1288	31	18	319.7	1368.4
1359	28	19	158.1	160.9
1360	29	19	97.3	781.6
1361	30	19	184.2	325.8
1364	33	19	264.1	339.9
1416	11	20	2.9	0.0
1428	23	20	6.9	113.1
1433	28	20	182.2	15.9
1434	29	20	1145.0	606.9
1435	30	20	2076.7	204.1
1436	31	20	1566.6	409.3
1437	32	20	184.6	286.4
1493	14	21	11.6	0.0
1494	15	21	1.5	0.0
1496	17	21	13.5	42.4
1498	19	21	0.0	413.9
1504	25	21	45.5	101.7
1505	26	21	259.1	253.8
1506	27	21	1314.2	73.8
1507	28	21	98.2	191.7
1508	29	21	1435.6	353.9
1509	30	21	1413.1	649.5
1510	31	21	1009.1	644.6
1511	32	21	37.8	467.0
1557	4	22	0.0	0.0
1558	5	22	0.0	0.0
1566	13	22	236.0	0.0
1568	15	22	0.0	274.7
1569	16	22	44.8	0.0
1574	21	22	0.6	163.4

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
1575	22	22	0.0	2246.4
1580	27	22	790.0	902.2
1581	28	22	1434.1	242.7
1582	29	22	724.3	298.0
1583	30	22	2536.3	484.7
1584	31	22	1130.1	538.1
1639	12	23	4.0	0.0
1640	13	23	5.9	0.0
1642	15	23	0.0	54.3
1646	19	23	50.7	0.0
1647	20	23	1.8	947.4
1648	21	23	5.0	0.0
1649	22	23	119.5	1346.1
1650	23	23	9.6	3585.8
1651	24	23	316.5	10735.0
1652	25	23	96.3	3928.6
1653	26	23	721.0	560.3
1654	27	23	332.5	4428.8
1655	28	23	913.7	556.6
1656	29	23	130.9	70.7
1657	30	23	1316.2	1532.8
1658	31	23	1367.5	1251.8
1659	32	23	0.0	345.5
1717	16	24	163.9	35.2
1718	17	24	91.0	0.0
1722	21	24	0.0	1279.3
1723	22	24	568.5	635.3
1724	23	24	711.5	1410.9
1725	24	24	173.0	3156.3
1726	25	24	189.0	1425.9
1727	26	24	408.4	3151.1
1728	27	24	100.0	6783.5
1729	28	24	866.0	1014.2
1730	29	24	236.0	449.6
1731	30	24	539.4	912.2
1732	31	24	122.8	1488.2
1786	11	25	0.0	56.7
1787	12	25	6.4	23.9

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
1789	14	25	97.0	0.0
1790	15	25	328.4	0.0
1794	19	25	769.0	1379.9
1795	20	25	669.5	527.7
1796	21	25	4160.0	2030.4
1797	22	25	1196.9	1816.7
1798	23	25	2762.1	951.5
1799	24	25	346.1	2475.7
1800	25	25	737.6	1728.5
1801	26	25	627.0	2273.4
1802	27	25	520.5	3947.8
1803	28	25	386.9	2278.6
1804	29	25	1296.0	1113.3
1805	30	25	1018.4	738.5
1806	31	25	893.3	3243.0
1809	34	25	89.1	679.0
1859	10	26	4.0	396.4
1862	13	26	0.0	127.0
1863	14	26	114.2	0.0
1864	15	26	44.6	0.0
1865	16	26	38.0	0.0
1868	19	26	662.0	698.6
1869	20	26	0.0	378.7
1870	21	26	3284.9	2624.9
1871	22	26	3178.7	2309.4
1872	23	26	945.8	3133.7
1873	24	26	214.4	5702.6
1874	25	26	860.0	2849.9
1875	26	26	4738.7	4470.7
1876	27	26	605.5	7229.7
1877	28	26	924.1	2531.4
1878	29	26	718.9	7251.8
1879	30	26	1289.1	1169.9
1880	31	26	1039.7	4254.5
1881	32	26	251.2	2847.3
1883	34	26	158.7	506.9
1932	9	27	1195.8	39.8
1933	10	27	42.3	518.5

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
1938	15	27	49.3	53.1
1941	18	27	236.3	0.0
1942	19	27	1790.8	2282.6
1943	20	27	533.3	1986.0
1944	21	27	85.4	2138.3
1945	22	27	2078.7	3749.2
1946	23	27	4156.3	1455.1
1947	24	27	3751.5	3456.8
1948	25	27	4289.8	1913.3
1949	26	27	4645.7	2603.1
1950	27	27	1471.2	2138.4
1951	28	27	78.4	3763.8
1952	29	27	2212.5	8880.6
1953	30	27	2343.2	6404.7
1954	31	27	378.7	9203.5
1955	32	27	134.3	8981.7
1956	33	27	233.1	5407.3
2005	8	28	189.1	0.0
2006	9	28	645.1	0.0
2007	10	28	134.4	103.2
2008	11	28	28.1	130.9
2009	12	28	0.0	255.0
2011	14	28	53.0	0.0
2012	15	28	101.0	0.0
2013	16	28	69.0	392.3
2014	17	28	267.2	0.0
2015	18	28	1400.5	460.0
2016	19	28	1651.3	655.6
2017	20	28	4664.1	2799.8
2018	21	28	1099.9	2478.9
2019	22	28	1223.4	3815.6
2020	23	28	7836.3	2688.6
2021	24	28	6506.3	969.8
2022	25	28	11117.1	3783.0
2023	26	28	282.9	906.6
2024	27	28	517.8	4984.4
2025	28	28	1017.4	3953.7
2026	29	28	2301.5	6456.9

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
2027	30	28	3702.9	7608.2
2028	31	28	5997.8	11717.2
2029	32	28	346.7	8808.0
2030	33	28	0.2	3960.4
2072	1	29	6.0	84.3
2073	2	29	306.5	76.7
2080	9	29	1140.9	45.3
2081	10	29	198.9	58.1
2082	11	29	501.9	126.7
2083	12	29	215.1	151.5
2084	13	29	1.6	0.0
2086	15	29	6.6	0.0
2087	16	29	331.9	1022.2
2088	17	29	501.2	481.0
2089	18	29	2470.1	715.4
2090	19	29	1289.8	469.5
2091	20	29	4717.3	5356.6
2092	21	29	6081.3	2748.3
2093	22	29	2931.3	4112.8
2094	23	29	10950.6	3991.5
2095	24	29	10516.9	682.5
2096	25	29	9851.4	3044.7
2097	26	29	11244.1	4209.1
2098	27	29	3795.4	3191.7
2099	28	29	3399.7	7569.1
2100	29	29	343.3	10756.4
2101	30	29	5273.5	13484.9
2102	31	29	883.7	9044.4
2103	32	29	1626.7	2307.1
2104	33	29	2044.2	3365.9
2147	2	30	4.6	81.2
2149	4	30	21.6	4.8
2154	9	30	1308.8	23.5
2155	10	30	452.0	0.0
2156	11	30	1127.7	0.0
2158	13	30	153.1	1.1
2159	14	30	0.0	0.0
2160	15	30	156.3	0.0

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
2162	17	30	0.0	0.0
2163	18	30	707.3	809.7
2165	20	30	4198.1	1193.5
2166	21	30	3190.9	100.4
2167	22	30	6050.0	1484.3
2168	23	30	10443.4	2879.2
2169	24	30	4113.9	325.2
2170	25	30	12274.3	166.1
2171	26	30	8056.1	6367.8
2172	27	30	5743.1	4075.0
2173	28	30	12153.7	6470.0
2174	29	30	2338.1	27955.3
2175	30	30	1407.1	9713.2
2176	31	30	1674.5	10613.9
2177	32	30	1194.1	8609.9
2178	33	30	448.5	4355.4
2179	34	30	47.0	6381.3
2225	6	31	0.1	0.0
2227	8	31	0.0	0.0
2228	9	31	37.0	0.0
2229	10	31	68.0	11.3
2230	11	31	268.7	0.0
2233	14	31	21.8	0.0
2234	15	31	907.9	0.0
2235	16	31	0.0	0.0
2236	17	31	548.3	0.0
2237	18	31	3068.9	106.0
2238	19	31	3518.0	227.8
2239	20	31	5438.2	1486.9
2240	21	31	6443.7	1575.9
2241	22	31	3540.5	3168.0
2242	23	31	2031.5	4294.4
2243	24	31	2416.6	1183.7
2244	25	31	3892.7	6007.8
2245	26	31	2331.2	4021.6
2246	27	31	8580.8	4359.3
2247	28	31	2347.7	18740.9
2248	29	31	5481.2	10386.7

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
2249	30	31	3947.2	7243.5
2250	31	31	7040.0	6549.5
2251	32	31	6828.9	8177.6
2252	33	31	1296.6	3025.2
2294	1	32	403.8	0.0
2296	3	32	6.3	10.3
2297	4	32	0.1	0.0
2299	6	32	0.1	0.0
2301	8	32	59.8	0.0
2302	9	32	0.1	0.0
2303	10	32	155.0	13.0
2304	11	32	4.8	0.0
2305	12	32	0.0	0.0
2307	14	32	0.9	0.0
2308	15	32	1827.9	66.8
2309	16	32	102.1	23.4
2310	17	32	440.2	1158.7
2311	18	32	738.0	715.1
2312	19	32	2573.1	299.2
2313	20	32	3076.9	440.6
2314	21	32	3832.4	282.4
2315	22	32	2793.7	1002.1
2316	23	32	407.4	7510.2
2317	24	32	5337.9	9125.4
2318	25	32	1791.6	9807.7
2319	26	32	11580.8	963.8
2320	27	32	7233.2	6024.2
2321	28	32	7457.3	2814.8
2322	29	32	22350.9	2053.2
2323	30	32	17141.5	3930.0
2324	31	32	12437.6	4664.1
2325	32	32	9079.4	3891.9
2326	33	32	1170.9	14056.5
2328	35	32	511.8	4873.1
2368	1	33	308.4	0.0
2374	7	33	10.7	0.0
2379	12	33	3.8	0.0
2380	13	33	30.9	0.0

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
2381	14	33	94.1	0.0
2382	15	33	385.5	45.4
2383	16	33	536.5	537.1
2384	17	33	2705.4	491.4
2385	18	33	663.1	683.3
2386	19	33	3313.3	1276.1
2387	20	33	1099.9	455.6
2388	21	33	4377.9	391.2
2389	22	33	1600.2	2394.4
2390	23	33	1109.2	4257.7
2391	24	33	3169.3	3887.8
2392	25	33	9825.8	15217.1
2393	26	33	10188.9	1479.8
2394	27	33	19623.3	3048.3
2395	28	33	24398.6	1986.3
2396	29	33	18065.7	13119.0
2397	30	33	10846.1	4231.2
2398	31	33	2130.4	9100.1
2399	32	33	8869.2	20822.6
2400	33	33	7529.0	11556.7
2401	34	33	930.6	5500.7
2402	35	33	960.1	2368.0
2448	7	34	0.0	0.0
2449	8	34	0.0	0.0
2454	13	34	2.4	0.0
2456	15	34	3.1	0.0
2458	17	34	617.7	840.8
2459	18	34	1126.2	3526.4
2460	19	34	2145.8	202.6
2461	20	34	299.0	564.3
2462	21	34	1983.4	67.7
2463	22	34	1067.2	2281.4
2465	24	34	7985.2	1465.6
2466	25	34	15666.4	6230.2
2467	26	34	11160.2	10074.6
2468	27	34	13431.8	11040.7
2469	28	34	19695.2	5769.1
2470	29	34	9455.2	3252.7

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
2471	30	34	17114.2	5371.1
2472	31	34	10119.7	8277.9
2473	32	34	7132.3	8562.8
2474	33	34	10871.6	7375.0
2475	34	34	2150.6	11447.6
2518	3	35	15.8	0.0
2519	4	35	28.2	0.0
2523	8	35	141.3	0.0
2528	13	35	40.5	0.0
2529	14	35	159.2	172.9
2530	15	35	0.8	62.1
2531	16	35	1418.9	92.7
2532	17	35	599.7	175.9
2533	18	35	312.4	1192.0
2534	19	35	72.4	1996.6
2535	20	35	1123.0	1807.3
2536	21	35	1385.4	68.1
2537	22	35	4036.4	4417.0
2538	23	35	6472.8	4278.0
2539	24	35	2105.1	1728.6
2540	25	35	3805.6	2645.9
2541	26	35	5531.2	2398.5
2542	27	35	6500.3	6492.0
2543	28	35	14895.6	4700.0
2544	29	35	8588.9	1867.6
2545	30	35	14354.0	3416.3
2546	31	35	17553.3	11781.1
2548	33	35	4338.4	3880.7
2549	34	35	4569.4	5678.8
2593	4	36	6.0	0.0
2599	10	36	127.0	0.0
2603	14	36	195.4	105.4
2604	15	36	0.0	11.3
2605	16	36	1034.2	148.5
2606	17	36	470.4	382.2
2607	18	36	418.9	140.4
2608	19	36	1301.6	0.0
2609	20	36	492.9	136.2

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
2610	21	36	1244.5	350.8
2611	22	36	3773.7	2078.0
2612	23	36	7047.2	6389.2
2613	24	36	5844.5	4352.1
2614	25	36	3434.3	2364.1
2615	26	36	2753.7	2391.5
2616	27	36	1365.5	13083.2
2617	28	36	9295.9	6755.4
2618	29	36	11080.9	5039.0
2619	30	36	13591.7	9985.5
2620	31	36	7153.1	1454.7
2622	33	36	5220.9	3618.8
2623	34	36	0.0	5260.0
2666	3	37	0.0	0.0
2669	6	37	0.0	0.0
2673	10	37	0.0	0.0
2674	11	37	135.1	0.0
2676	13	37	16.3	0.0
2677	14	37	61.2	0.0
2678	15	37	253.5	107.3
2679	16	37	1766.9	0.0
2680	17	37	474.9	51.3
2681	18	37	562.6	61.4
2682	19	37	771.9	3593.2
2684	21	37	1982.1	208.2
2685	22	37	1983.1	278.2
2686	23	37	1107.6	939.4
2687	24	37	3588.4	5032.5
2688	25	37	1405.0	5338.1
2690	27	37	1643.2	4456.4
2691	28	37	8050.7	4241.4
2692	29	37	2718.4	7433.9
2743	6	38	0.0	0.0
2748	11	38	21.9	0.0
2749	12	38	147.9	0.0
2750	13	38	327.9	0.0
2751	14	38	1174.5	10.9
2752	15	38	254.3	0.0

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
2753	16	38	460.0	86.0
2754	17	38	1138.9	170.5
2755	18	38	2159.2	0.0
2756	19	38	361.2	83.3
2757	20	38	1832.4	20.2
2758	21	38	5248.8	0.0
2759	22	38	3011.9	2166.6
2760	23	38	2892.7	3762.1
2761	24	38	2450.0	739.5
2762	25	38	251.0	1602.7
2765	28	38	3645.4	12140.2
2766	29	38	1624.2	8721.1
2767	30	38	8100.4	8946.0
2768	31	38	456.0	4379.4
2816	5	39	75.2	0.0
2818	7	39	0.0	0.0
2821	10	39	0.0	46.0
2822	11	39	42.8	99.7
2824	13	39	921.1	0.0
2825	14	39	646.0	159.3
2826	15	39	490.4	48.7
2827	16	39	147.7	0.4
2829	18	39	732.9	0.0
2830	19	39	316.4	82.1
2831	20	39	875.0	1526.4
2832	21	39	1571.1	373.0
2833	22	39	3344.4	1122.1
2834	23	39	2343.5	2012.5
2835	24	39	489.9	1973.9
2837	26	39	0.0	5808.7
2839	28	39	2482.3	5.3
2841	30	39	5446.2	354.1
2842	31	39	382.3	9411.3
2843	32	39	5311.4	2570.0
2897	12	40	0.0	21.6
2898	13	40	166.7	0.0
2899	14	40	159.2	156.3
2900	15	40	406.2	799.8

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
2902	17	40	1075.1	62.3
2903	18	40	1870.4	156.7
2904	19	40	315.9	0.0
2905	20	40	501.6	10.2
2906	21	40	8496.1	39.6
2907	22	40	6366.2	113.9
2908	23	40	11035.9	338.5
2909	24	40	1078.4	2752.0
2910	25	40	1442.0	4269.4
2912	27	40	3521.0	5881.9
2913	28	40	4273.6	2027.0
2914	29	40	8562.9	260.1
2915	30	40	5489.9	0.0
2916	31	40	2178.1	1707.7
2962	3	41	3.8	0.0
2966	7	41	52.8	140.8
2971	12	41	0.2	326.4
2973	14	41	41.4	0.0
2974	15	41	376.5	42.7
2975	16	41	129.5	0.0
2976	17	41	82.3	52.5
2977	18	41	207.6	0.0
2978	19	41	182.3	153.3
2979	20	41	846.1	210.4
2980	21	41	1613.1	30.5
2981	22	41	1702.0	627.1
2982	23	41	4607.7	562.0
2984	25	41	4209.0	9109.6
2985	26	41	6733.3	9458.9
2986	27	41	3467.8	7916.4
2987	28	41	1332.5	7337.1
2989	30	41	1442.1	11.2
2990	31	41	1014.3	121.1
2991	32	41	7710.2	0.0
2992	33	41	7.7	0.0
3038	5	42	60.4	0.0
3040	7	42	188.7	0.0
3041	8	42	201.3	30.8

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
3043	10	42	8.9	1237.8
3045	12	42	0.0	177.2
3046	13	42	119.5	56.7
3047	14	42	50.0	191.5
3048	15	42	12.5	88.5
3050	17	42	4.3	54.6
3051	18	42	56.6	119.6
3052	19	42	2139.3	846.0
3053	20	42	802.8	157.1
3054	21	42	818.4	938.9
3055	22	42	3576.3	3823.5
3056	23	42	444.4	3186.2
3057	24	42	5942.1	7047.0
3058	25	42	531.1	8680.5
3059	26	42	3425.0	12467.1
3060	27	42	1437.0	7954.4
3061	28	42	326.0	10338.9
3062	29	42	520.1	14513.1
3067	34	42	409.5	0.0
3113	6	43	50.1	0.0
3117	10	43	56.0	525.7
3119	12	43	0.0	94.0
3120	13	43	135.5	0.0
3121	14	43	417.0	2.6
3123	16	43	16.8	0.0
3124	17	43	71.4	0.0
3125	18	43	0.0	246.0
3126	19	43	204.0	304.3
3127	20	43	202.1	152.4
3128	21	43	891.5	704.2
3129	22	43	2481.9	3491.7
3130	23	43	2669.6	1717.9
3131	24	43	11926.2	4692.2
3132	25	43	4664.0	3202.2
3133	26	43	2595.6	11896.9
3134	27	43	288.9	9962.0
3136	29	43	775.6	7622.1
3140	33	43	3528.0	0.0

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
3188	7	44	6.3	0.0
3189	8	44	4.6	0.0
3191	10	44	290.3	0.0
3192	11	44	75.9	14.8
3196	15	44	110.4	93.2
3198	17	44	131.1	0.0
3199	18	44	67.4	390.2
3200	19	44	168.5	326.2
3201	20	44	218.4	81.1
3202	21	44	651.3	55.9
3203	22	44	2367.0	641.0
3204	23	44	8898.0	2903.3
3205	24	44	16383.8	5358.9
3206	25	44	16873.7	1681.2
3207	26	44	9557.7	2691.7
3208	27	44	1173.7	7661.8
3213	32	44	1135.1	58.1
3214	33	44	3649.9	478.4
3215	34	44	3283.1	4775.3
3261	6	45	109.5	0.0
3262	7	45	128.8	0.0
3263	8	45	5601.1	0.0
3264	9	45	1022.0	43.7
3265	10	45	54.2	17.0
3266	11	45	393.2	200.7
3267	12	45	63.9	139.5
3268	13	45	354.9	284.3
3269	14	45	261.1	195.3
3270	15	45	95.5	297.3
3273	18	45	50.8	210.5
3274	19	45	65.4	184.0
3275	20	45	158.3	1183.7
3276	21	45	1390.8	1763.7
3277	22	45	5972.8	1863.1
3278	23	45	13050.6	7540.9
3279	24	45	18351.2	8897.8
3280	25	45	15661.6	8355.0
3281	26	45	17038.6	11859.2

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
3282	27	45	11500.5	7826.1
3283	28	45	1916.9	5267.1
3284	29	45	1052.0	3410.0
3288	33	45	2513.7	1545.6
3289	34	45	11664.4	487.9
3337	8	46	100.1	188.1
3338	9	46	0.0	44.5
3339	10	46	103.6	0.0
3340	11	46	54.2	137.4
3343	14	46	115.0	0.0
3344	15	46	3.0	10.6
3345	16	46	142.7	0.0
3346	17	46	9.8	0.0
3347	18	46	298.1	148.6
3348	19	46	136.6	471.2
3349	20	46	210.3	668.3
3350	21	46	1012.8	949.0
3351	22	46	5020.5	1312.5
3352	23	46	8299.3	7787.8
3353	24	46	13368.6	7452.3
3354	25	46	11861.2	8293.6
3355	26	46	12443.1	8180.1
3356	27	46	13419.0	9300.6
3357	28	46	4839.3	8640.2
3358	29	46	655.7	10648.4
3410	7	47	0.0	25.2
3412	9	47	0.0	0.0
3413	10	47	398.6	26.7
3414	11	47	241.9	205.5
3415	12	47	42.7	218.7
3416	13	47	274.8	44.0
3420	17	47	146.7	88.9
3421	18	47	176.5	272.6
3422	19	47	134.2	911.4
3423	20	47	47.6	445.3
3424	21	47	483.1	849.1
3425	22	47	3185.4	3064.2
3426	23	47	1810.5	2194.7

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
3427	24	47	3280.7	5007.7
3428	25	47	12040.9	10343.6
3429	26	47	12737.8	5301.5
3430	27	47	9893.3	6859.9
3431	28	47	3549.6	8700.8
3432	29	47	1948.9	9597.9
3433	30	47	1195.9	9841.2
3435	32	47	181.8	943.2
3485	8	48	38.1	137.2
3486	9	48	58.9	53.3
3487	10	48	965.3	190.8
3488	11	48	601.1	186.0
3489	12	48	37.7	306.7
3490	13	48	393.5	68.3
3492	15	48	63.2	11.7
3493	16	48	51.3	0.0
3494	17	48	126.6	0.0
3495	18	48	664.3	0.0
3496	19	48	809.2	447.6
3497	20	48	9.6	1663.5
3498	21	48	2255.1	1627.4
3499	22	48	7666.9	1083.8
3500	23	48	7331.3	3173.0
3501	24	48	9157.5	4120.8
3502	25	48	5814.3	2499.6
3503	26	48	12762.0	1604.0
3504	27	48	4933.1	2918.5
3505	28	48	5764.1	2731.0
3506	29	48	2415.8	4351.0
3507	30	48	383.1	7266.2
3508	31	48	1026.3	6861.3
3509	32	48	2799.3	5871.2
3559	8	49	191.2	0.0
3560	9	49	0.4	0.0
3561	10	49	874.4	15.9
3562	11	49	805.6	0.0
3563	12	49	176.6	30.1
3564	13	49	3.5	0.0

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
3568	17	49	348.7	24.9
3569	18	49	808.2	61.9
3570	19	49	541.1	120.2
3571	20	49	125.1	2309.9
3572	21	49	4142.3	894.8
3573	22	49	5902.1	1358.3
3574	23	49	9734.1	2327.5
3575	24	49	13701.8	4968.6
3576	25	49	9229.4	315.1
3577	26	49	15067.6	423.9
3578	27	49	7077.1	2626.9
3579	28	49	2794.3	2021.9
3580	29	49	6470.2	2751.8
3581	30	49	592.7	2060.7
3582	31	49	2177.2	1587.1
3583	32	49	4383.4	6391.8
3633	8	50	54.6	0.0
3634	9	50	603.8	0.0
3635	10	50	572.4	0.0
3636	11	50	39.9	0.0
3637	12	50	3.0	0.7
3640	15	50	10.2	0.0
3641	16	50	448.3	0.0
3642	17	50	48.5	55.8
3644	19	50	194.8	86.7
3645	20	50	910.8	2753.3
3646	21	50	1709.6	3186.0
3647	22	50	2000.1	314.1
3648	23	50	7532.0	536.3
3649	24	50	5743.9	734.0
3650	25	50	6783.5	147.8
3651	26	50	10242.0	976.0
3652	27	50	14456.4	4403.5
3653	28	50	6270.7	2432.1
3654	29	50	3794.8	2604.8
3655	30	50	2491.2	2588.4
3656	31	50	1358.8	1852.2
3657	32	50	1867.2	465.8

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
3658	33	50	1159.4	115.2
3659	34	50	56.4	0.0
3707	8	51	0.0	0.0
3708	9	51	14.4	0.0
3709	10	51	141.7	0.0
3710	11	51	377.4	0.0
3711	12	51	15.2	0.0
3712	13	51	59.0	0.0
3713	14	51	105.5	0.0
3717	18	51	116.8	0.0
3718	19	51	727.3	857.0
3719	20	51	604.6	2600.7
3720	21	51	7559.1	3112.7
3721	22	51	1943.0	175.2
3722	23	51	2151.2	361.0
3723	24	51	1806.4	159.7
3724	25	51	4102.1	3.8
3725	26	51	4789.0	147.1
3726	27	51	5017.3	490.3
3727	28	51	9060.0	1224.8
3728	29	51	2886.2	1710.1
3729	30	51	1807.2	1375.1
3730	31	51	466.8	281.3
3731	32	51	2830.8	0.0
3780	7	52	0.0	0.0
3783	10	52	358.4	0.0
3786	13	52	60.0	0.0
3787	14	52	57.3	0.0
3789	16	52	0.0	6.0
3791	18	52	85.8	168.7
3792	19	52	419.6	1303.9
3793	20	52	1857.3	1844.7
3794	21	52	3218.1	107.9
3795	22	52	5056.4	238.6
3796	23	52	544.5	289.8
3797	24	52	1117.9	376.3
3798	25	52	2840.4	1138.0
3799	26	52	3790.5	742.8

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
3800	27	52	3468.5	99.4
3801	28	52	4071.8	744.2
3802	29	52	1750.1	766.4
3803	30	52	4255.1	62.1
3804	31	52	225.0	0.0
3805	32	52	217.1	0.0
3855	8	53	0.0	0.0
3864	17	53	59.5	161.2
3865	18	53	4221.8	423.8
3866	19	53	2982.4	316.8
3867	20	53	198.9	492.9
3868	21	53	246.3	177.7
3869	22	53	948.5	931.3
3870	23	53	148.7	0.0
3871	24	53	261.6	686.5
3872	25	53	1834.7	318.3
3873	26	53	1830.3	326.2
3874	27	53	2862.6	1236.7
3875	28	53	1657.2	2156.8
3876	29	53	2641.8	37.2
3877	30	53	1254.7	0.0
3878	31	53	473.3	0.0
3936	15	54	3.1	0.0
3938	17	54	0.1	48.6
3939	18	54	1091.5	183.3
3940	19	54	1114.5	845.6
3941	20	54	202.4	1397.5
3944	23	54	10.9	1094.8
3945	24	54	390.9	793.7
3946	25	54	1108.8	812.2
3947	26	54	575.8	3622.5
3948	27	54	1788.2	1103.0
3949	28	54	913.4	66.4
3950	29	54	298.6	0.0
4000	5	55	0.0	0.0
4004	9	55	0.0	4.2
4008	13	55	0.0	0.0
4012	17	55	249.9	151.8

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
4014	19	55	135.9	0.0
4018	23	55	78.0	787.9
4019	24	55	4.6	590.7
4020	25	55	34.0	120.8
4021	26	55	214.3	2371.2
4022	27	55	510.3	776.1
4023	28	55	243.5	117.4
4024	29	55	32.7	0.0
4081	12	56	11.2	0.0
4085	16	56	0.1	0.0
4087	18	56	8.7	237.3
4088	19	56	14.3	1127.2
4092	23	56	46.4	392.9
4093	24	56	10.0	1806.7
4094	25	56	602.4	1301.3
4095	26	56	971.4	2507.1
4096	27	56	219.3	1519.6
4158	15	57	0.1	0.0
4159	16	57	0.0	0.0
4160	17	57	36.1	46.7
4162	19	57	2.8	3091.5
4163	20	57	214.0	3324.5
4164	21	57	74.9	2153.5
4165	22	57	21.2	1811.2
4166	23	57	23.2	2052.0
4167	24	57	1034.1	1135.8
4168	25	57	1050.0	5486.1
4169	26	57	410.9	8869.7
4228	11	58	11.2	0.0
4229	12	58	0.0	0.0
4235	18	58	5.7	2049.4
4236	19	58	28.6	2294.4
4237	20	58	160.5	3918.1
4238	21	58	64.4	1181.5
4240	23	58	0.0	450.3
4241	24	58	451.8	3907.9
4242	25	58	200.8	2339.7
4302	11	59	41.4	0.0

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
4306	15	59	118.1	415.0
4307	16	59	54.9	256.7
4311	20	59	116.3	118.9
4313	22	59	18.8	70.9
4314	23	59	2.1	1327.3
4315	24	59	87.1	2445.4
4376	11	60	0.0	3.4
4382	17	60	0.0	200.0
4383	18	60	13.4	1202.3
4384	19	60	100.0	95.5
4385	20	60	0.0	122.8
4386	21	60	0.0	241.6
4387	22	60	0.0	3317.8
4388	23	60	0.0	3667.0
4389	24	60	3.7	957.2
4442	3	61	0.7	0.0
4443	4	61	0.0	0.0
4458	19	61	0.0	0.0
4459	20	61	0.0	30.1
4460	21	61	0.0	58.0
4461	22	61	0.0	2149.7
4462	23	61	0.0	332.0
4463	24	61	0.0	780.2
4516	3	62	0.0	0.0
4522	9	62	0.0	0.0
4527	14	62	0.0	6.6
4528	15	62	0.0	3.5
4529	16	62	0.0	0.0
4531	18	62	0.0	0.0
4532	19	62	0.0	0.0
4533	20	62	0.0	0.0
4534	21	62	0.0	147.7
4535	22	62	0.0	562.9
4536	23	62	5.1	260.3
4537	24	62	0.0	291.4
4543	30	62	0.0	0.0
4590	3	63	0.0	0.0
4594	7	63	0.0	0.0

Table E-1 (continued).

Grid Cell ID^a	i^b	j^b	2009 Natural Gas Production (MMscf/y)^c	2013 Natural Gas Production (MMscf/y)^d
4595	8	63	0.0	0.0
4597	10	63	0.0	0.0
4599	12	63	0.0	0.0
4600	13	63	0.0	123.8
4605	18	63	0.0	0.0
4606	19	63	0.0	0.0
4607	20	63	0.0	0.0
4608	21	63	0.0	136.0
4609	22	63	424.6	83.6
4610	23	63	0.0	79.8
4672	11	64	0.0	0.0
4674	13	64	0.0	0.0
4679	18	64	0.0	0.0
4680	19	64	0.0	0.0
4682	21	64	0.0	0.0
4683	22	64	15.6	0.0
4684	23	64	64.2	0.0
4753	18	65	0.0	0.0
4756	21	65	0.0	0.0
4757	22	65	0.0	0.0
4758	23	65	0.0	0.0

(a) Grid ID represents the number of the cell from the domain. Starting at the lower left corner; followed by cell values in row-major order.

(b) i,j represent the coordinates of each grid cell in the domain, with $x_0=140$, $y_0=-680$ (4km by 4km domain. [6])

(c) Represents sum of Natural Gas Production for all the sites in the special emission inventory, within each grid cell.

(d) Obtained from Texas Rail Road Commission.

For each grid cell, Table E-2 shows final data set of estimated emissions for 2009.

Table E-2. Estimated methane, ethane and propane emissions, as well as methane to ethane and methane to propane ratios, for each grid cell, for 2009.

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
1066	1.8	0.2	0.2	14.5	28.0
1139	0.1	0.0	0.0	14.4	27.7
1285	0.1	0.0	0.0	14.0	25.8
1287	0.0	0.0	0.0	14.8	29.2
1288	2.2	0.3	0.2	15.1	30.9
1359	0.1	0.0	0.0	13.2	26.3
1360	1.8	0.2	0.2	14.2	27.3
1361	0.7	0.1	0.1	15.2	33.6
1364	0.0	0.0	0.0	15.6	33.5
1416	0.5	0.1	0.1	11.6	18.4
1428	1.4	0.2	0.2	12.5	20.3
1433	1.0	0.1	0.1	12.8	21.9
1434	2.9	0.4	0.4	12.0	19.7
1435	2.9	0.4	0.2	15.3	38.8
1436	3.5	0.4	0.1	18.8	67.9
1437	0.1	0.0	0.0	17.4	45.6
1493	0.9	0.1	0.1	11.4	18.0
1494	1.4	0.2	0.2	11.6	18.8
1496	1.5	0.2	0.2	11.4	18.1
1498	0.3	0.0	0.0	17.9	57.9
1504	2.0	0.3	0.3	12.5	20.3
1505	5.7	0.9	0.9	11.5	17.4
1506	3.8	0.7	0.6	10.5	16.7
1507	0.7	0.1	0.1	12.0	20.4
1508	7.1	1.0	0.9	12.7	21.3
1509	1.5	0.2	0.1	14.9	34.4
1510	10.0	1.2	0.4	16.0	73.3
1511	0.3	0.0	0.0	18.5	56.8
1557	0.6	0.1	0.1	11.8	18.2
1558	0.6	0.1	0.1	11.7	17.8
1566	0.9	0.4	0.6	4.2	4.1
1568	0.0	0.0	0.0	NA	NA
1569	2.8	0.5	0.4	11.5	18.3

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
1574	1.4	0.2	0.2	11.6	18.4
1575	1.4	0.2	0.2	12.0	19.0
1580	4.7	0.8	0.8	11.0	16.7
1581	3.4	0.5	0.4	12.3	20.7
1582	7.2	0.9	0.6	15.1	31.7
1583	10.9	1.1	0.5	18.1	59.3
1584	3.5	0.4	0.2	17.0	50.9
1639	0.0	0.0	0.0	5.4	5.6
1640	0.0	0.0	0.0	7.8	9.2
1642	0.7	0.1	0.1	11.9	18.5
1646	0.1	0.0	0.0	12.4	23.4
1647	0.2	0.0	0.0	14.3	31.5
1648	0.7	0.1	0.1	11.6	19.2
1649	3.3	0.5	0.5	12.0	19.4
1650	4.2	0.6	0.5	12.6	21.0
1651	1.0	0.1	0.1	15.0	34.5
1652	3.1	0.5	0.4	12.4	20.3
1653	9.8	1.6	1.5	11.7	18.1
1654	3.3	0.5	0.4	12.7	20.9
1655	4.6	0.7	0.7	11.5	17.7
1656	1.6	0.2	0.2	13.1	22.3
1657	2.6	0.3	0.1	17.0	52.5
1658	13.2	1.2	0.3	21.0	109.7
1659	0.0	0.0	0.0	16.8	42.4
1717	5.4	0.9	0.8	11.3	18.4
1718	5.5	1.0	0.9	10.5	15.9
1722	0.0	0.0	0.0	11.5	17.3
1723	8.6	1.3	1.2	12.1	19.5
1724	2.3	0.4	0.3	11.9	19.4
1725	0.0	0.0	0.0	13.6	24.2
1726	0.3	0.0	0.0	13.2	23.3
1727	7.2	1.1	1.0	12.0	18.9
1728	3.4	0.5	0.5	11.8	18.5
1729	1.8	0.3	0.2	12.9	22.6
1730	0.0	0.0	0.0	14.6	28.4
1731	0.1	0.0	0.0	16.5	38.8
1732	0.1	0.0	0.0	17.6	47.8

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
1786	0.0	0.0	0.0	6.4	7.0
1787	1.8	0.5	0.4	7.0	12.0
1789	0.5	0.1	0.1	12.8	27.9
1790	6.6	1.0	0.8	12.3	23.3
1794	6.2	1.1	1.0	10.9	17.8
1795	4.9	0.9	0.8	10.0	16.2
1796	25.2	4.7	4.7	10.1	14.8
1797	5.8	1.0	0.9	11.1	16.8
1798	2.2	0.4	0.3	11.9	22.4
1799	2.2	0.3	0.3	13.3	23.6
1800	0.3	0.0	0.0	13.4	33.0
1801	0.2	0.0	0.0	14.8	32.9
1802	0.6	0.1	0.0	15.4	33.9
1803	2.3	0.3	0.2	14.6	28.9
1804	0.2	0.0	0.0	15.8	34.5
1805	0.3	0.0	0.0	17.1	43.2
1806	2.6	0.2	0.1	19.8	84.7
1809	0.1	0.0	0.0	16.9	41.7
1859	0.0	0.0	0.0	5.1	5.3
1862	0.0	0.0	0.0	NA	NA
1863	5.6	0.7	0.5	14.6	29.0
1864	2.3	0.2	0.1	18.8	68.8
1865	0.8	0.1	0.1	14.3	33.1
1868	32.5	5.2	4.4	11.6	20.1
1869	1.0	0.2	0.2	10.4	15.7
1870	7.9	1.6	1.7	9.4	12.6
1871	3.4	0.8	0.9	8.1	10.5
1872	0.6	0.1	0.1	11.5	18.2
1873	0.2	0.0	0.0	13.0	24.3
1874	0.2	0.0	0.0	15.0	33.9
1875	1.2	0.1	0.1	15.9	37.7
1876	0.1	0.0	0.0	15.4	32.5
1877	0.5	0.1	0.0	16.4	42.4
1878	0.2	0.0	0.0	17.0	42.3
1879	2.1	0.2	0.1	19.5	72.7
1880	3.8	0.4	0.2	18.4	60.3
1881	2.4	0.3	0.2	16.4	41.9

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
1883	0.2	0.0	0.0	17.3	44.9
1932	3.1	1.0	1.4	5.7	6.1
1933	0.2	0.1	0.1	8.2	10.2
1938	2.2	0.3	0.1	16.3	42.6
1941	2.5	0.5	0.4	10.0	17.2
1942	15.9	2.8	2.8	10.6	15.9
1943	7.4	1.3	1.3	10.5	15.6
1944	7.4	1.3	1.3	10.9	15.8
1945	6.8	1.6	1.9	8.0	9.9
1946	2.9	0.5	0.6	10.1	14.0
1947	2.1	0.3	0.2	15.3	33.7
1948	0.2	0.0	0.0	15.0	30.1
1949	7.8	0.8	0.4	18.0	51.6
1950	0.2	0.0	0.0	16.0	35.7
1951	0.0	0.0	0.0	16.6	39.8
1952	1.1	0.1	0.1	16.9	44.0
1953	0.2	0.0	0.0	18.2	53.8
1954	18.3	1.9	0.6	17.8	79.3
1955	2.5	0.3	0.1	17.7	48.7
1956	2.7	0.3	0.1	17.6	51.4
2005	2.2	0.4	0.4	10.9	16.1
2006	2.6	0.9	1.2	5.5	5.9
2007	1.4	0.3	0.3	9.1	11.9
2008	0.6	0.1	0.1	9.4	12.3
2009	0.0	0.0	0.0	NA	NA
2011	0.6	0.1	0.1	12.2	24.0
2012	1.2	0.2	0.1	13.6	31.1
2013	7.0	1.0	0.8	13.6	24.4
2014	7.6	1.1	1.0	12.3	20.4
2015	13.6	2.4	2.4	10.7	15.9
2016	24.1	4.3	4.3	10.5	15.3
2017	34.2	6.1	6.1	10.6	15.5
2018	12.0	2.3	2.4	9.8	13.8
2019	7.5	2.3	2.9	6.2	7.1
2020	3.5	0.8	0.9	8.2	10.6
2021	0.8	0.1	0.1	13.5	25.1
2022	3.9	0.4	0.2	17.8	51.0

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
2023	0.0	0.0	0.0	15.8	34.5
2024	0.7	0.1	0.0	16.7	42.6
2025	12.8	1.6	0.5	14.6	65.9
2026	0.3	0.0	0.0	18.0	51.7
2027	0.4	0.0	0.0	17.5	50.2
2028	47.2	4.6	2.0	19.0	63.4
2029	6.5	0.5	0.1	23.3	252.5
2030	0.1	0.0	0.0	19.0	63.1
2072	0.6	0.1	0.1	11.5	18.0
2073	3.5	0.6	0.5	10.7	17.5
2080	4.7	1.6	2.3	5.4	5.7
2081	0.8	0.2	0.3	7.3	8.7
2082	2.2	0.6	0.8	6.4	7.4
2083	1.7	0.4	0.5	7.3	8.9
2084	0.0	0.0	0.0	10.5	14.7
2086	0.6	0.1	0.1	14.9	33.5
2087	6.2	0.8	0.6	14.1	27.4
2088	12.4	2.1	2.0	11.0	16.8
2089	17.9	3.2	3.2	10.4	15.6
2090	8.4	1.5	1.5	10.4	15.1
2091	34.4	6.1	6.3	10.5	15.0
2092	18.4	3.9	4.3	8.7	11.7
2093	2.4	1.1	1.5	4.1	4.4
2094	17.4	4.8	4.0	6.7	11.9
2095	1.1	0.2	0.1	12.5	20.4
2096	0.2	0.0	0.0	14.8	29.2
2097	0.8	0.1	0.1	15.9	37.7
2098	6.0	0.6	0.3	18.2	54.5
2099	1.8	0.2	0.1	17.1	70.0
2100	0.4	0.0	0.0	19.0	63.3
2101	3.4	0.3	0.1	19.7	76.2
2102	1.3	0.1	0.0	19.7	79.1
2103	7.7	0.7	0.2	20.3	90.2
2104	2.5	0.2	0.1	19.8	78.7
2147	0.6	0.1	0.1	11.1	17.3
2149	0.0	0.0	0.0	NA	NA
2154	5.5	1.6	2.2	6.3	7.1

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
2155	0.9	0.2	0.2	8.5	12.7
2156	5.2	1.4	1.9	6.8	7.6
2158	4.7	0.8	0.7	11.3	17.4
2159	1.0	0.2	0.2	11.0	15.9
2160	1.9	0.4	0.4	9.6	13.0
2162	0.9	0.1	0.1	11.6	17.5
2163	8.3	1.4	1.4	11.1	16.5
2165	25.0	4.4	4.4	10.7	15.6
2166	13.4	3.1	3.7	8.1	10.0
2167	12.0	2.7	2.9	8.5	11.3
2168	6.4	0.8	0.6	14.4	28.6
2169	0.6	0.1	0.1	13.6	24.4
2170	0.7	0.1	0.0	16.2	38.6
2171	14.0	1.2	0.2	22.5	188.6
2172	4.1	0.4	0.1	20.3	106.7
2173	51.3	4.0	0.4	23.8	374.1
2174	1.8	0.2	0.1	19.9	87.8
2175	6.3	0.6	0.2	19.9	90.3
2176	16.1	1.5	0.5	20.7	95.8
2177	4.5	0.4	0.1	20.7	99.7
2178	17.8	1.6	0.6	20.3	86.3
2179	2.4	0.2	0.1	19.2	72.5
2225	0.1	0.0	0.0	10.3	14.4
2227	0.0	0.0	0.0	NA	NA
2228	1.5	0.3	0.3	9.6	12.6
2229	0.2	0.0	0.0	9.5	13.5
2230	1.5	0.4	0.5	7.5	9.1
2233	0.3	0.0	0.0	13.2	23.4
2234	5.1	1.1	1.3	8.5	11.2
2235	0.1	0.0	0.0	11.0	16.0
2236	6.2	1.0	1.0	11.4	17.1
2237	8.2	1.3	1.3	11.6	17.6
2238	21.5	3.7	3.7	10.8	16.1
2239	56.7	9.7	9.7	11.0	16.0
2240	32.7	5.5	5.2	11.1	17.3
2241	7.7	1.5	1.5	9.7	13.9
2242	8.2	1.4	0.8	11.3	26.7

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
2243	6.1	0.7	0.5	15.8	36.8
2244	6.8	0.7	0.4	17.5	52.4
2245	114.2	8.6	0.3	24.7	1001.0
2246	14.3	1.4	0.5	19.6	77.7
2247	1.4	0.1	0.0	19.6	88.8
2248	21.7	2.3	0.5	17.3	121.8
2249	7.6	0.7	0.1	21.7	139.9
2250	12.0	1.1	0.3	20.9	102.9
2251	163.6	12.9	0.7	23.8	651.2
2252	4.1	0.4	0.1	20.9	117.6
2294	7.9	1.3	1.1	11.3	20.2
2296	1.1	0.2	0.2	10.2	14.6
2297	0.9	0.3	0.4	5.0	5.8
2299	0.1	0.0	0.0	10.6	15.1
2301	0.9	0.2	0.2	10.6	14.9
2302	0.1	0.0	0.0	10.6	14.9
2303	1.1	0.2	0.2	10.4	14.8
2304	1.1	0.2	0.2	10.6	15.1
2305	0.5	0.1	0.1	11.3	16.7
2307	0.1	0.0	0.0	11.2	19.4
2308	4.5	1.3	1.7	6.2	7.0
2309	0.2	0.0	0.0	11.5	18.5
2310	5.1	0.7	0.6	13.4	23.5
2311	1.0	0.1	0.1	12.7	21.3
2312	7.7	1.2	1.2	11.7	18.1
2313	17.1	2.9	3.0	10.9	15.9
2314	17.3	3.1	3.2	10.3	14.8
2315	4.6	1.4	1.8	6.1	6.9
2316	2.0	0.3	0.3	12.0	20.7
2317	6.4	0.7	0.4	16.5	41.9
2318	1.4	0.1	0.1	18.3	56.1
2319	2.2	0.2	0.1	18.3	57.9
2320	14.3	1.2	0.3	21.6	130.1
2321	29.0	2.5	0.5	21.9	150.5
2322	145.9	11.5	1.1	23.8	352.7
2323	34.1	3.0	0.9	21.0	106.4
2324	66.5	5.3	0.7	23.3	262.9

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
2325	49.0	4.0	0.7	22.7	197.2
2326	54.0	4.2	0.3	24.2	479.5
2328	2.4	0.2	0.1	20.0	81.4
2368	2.7	0.5	0.4	10.8	18.9
2374	0.0	0.0	0.0	11.2	16.5
2379	0.0	0.0	0.0	12.1	19.0
2380	0.3	0.1	0.0	10.0	21.4
2381	1.3	0.2	0.1	14.2	30.7
2382	6.1	1.0	0.9	11.5	18.4
2383	4.9	0.7	0.6	12.6	21.7
2384	45.7	6.4	5.3	13.4	23.6
2385	6.1	0.8	0.5	14.4	31.2
2386	2.0	0.4	0.3	10.6	20.6
2387	6.0	1.1	1.0	10.6	16.4
2388	10.8	2.4	2.7	8.4	10.8
2389	11.1	2.8	2.7	7.4	11.2
2390	4.2	0.6	0.5	12.7	22.1
2391	6.9	0.7	0.3	18.8	67.6
2392	55.6	4.7	1.0	22.1	146.6
2393	5.0	0.5	0.2	18.2	64.7
2394	11.2	1.0	0.4	20.0	84.6
2395	37.0	3.3	0.9	21.1	113.7
2396	14.2	1.4	0.5	19.4	79.9
2397	7.1	0.6	0.2	20.5	97.8
2398	7.2	0.6	0.2	21.2	122.6
2399	118.7	9.3	0.8	23.9	397.1
2400	73.3	6.3	1.0	21.7	199.3
2401	2.1	0.2	0.0	21.3	117.7
2402	0.0	0.0	0.0	20.0	78.7
2448	0.0	0.0	0.0	11.6	17.5
2449	0.9	0.1	0.1	11.5	17.9
2454	0.0	0.0	0.0	13.2	22.6
2456	0.1	0.0	0.0	13.2	26.2
2458	1.7	0.2	0.2	13.3	27.8
2459	5.8	0.8	0.6	13.6	26.8
2460	8.8	1.6	1.6	10.6	15.3
2461	2.9	0.5	0.4	11.7	18.4

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
2462	2.9	0.7	0.7	8.3	11.3
2463	4.2	0.9	0.9	9.0	12.3
2465	6.3	0.8	0.6	14.8	31.4
2466	4.0	0.5	0.2	14.7	46.4
2467	0.9	0.1	0.1	17.4	45.9
2468	0.2	0.0	0.0	18.8	59.7
2469	7.9	0.8	0.3	19.6	82.8
2470	9.4	0.9	0.3	20.4	92.4
2471	12.3	1.2	0.4	19.9	81.3
2472	9.4	0.9	0.3	20.1	101.9
2473	69.5	5.6	0.8	23.1	242.2
2474	34.1	2.7	0.3	23.8	360.0
2475	0.3	0.0	0.0	20.9	98.0
2518	0.1	0.0	0.0	6.4	8.2
2519	0.2	0.1	0.1	6.9	8.3
2523	0.6	0.1	0.1	12.0	18.6
2528	0.2	0.0	0.0	11.5	25.4
2529	1.2	0.2	0.1	14.8	29.9
2530	0.2	0.0	0.0	14.6	32.9
2531	4.0	0.5	0.3	14.8	31.6
2532	10.9	1.0	0.3	20.1	96.1
2533	2.8	0.4	0.3	13.5	24.1
2534	9.7	1.9	1.2	9.7	22.1
2535	4.3	0.6	0.4	14.6	31.6
2536	8.5	1.2	1.0	13.1	23.0
2537	17.2	2.4	1.9	13.6	25.2
2538	9.6	1.3	1.0	14.1	27.3
2539	3.9	0.6	0.5	12.9	23.5
2540	2.4	0.3	0.2	15.8	34.8
2541	3.3	0.5	0.4	12.2	21.9
2542	0.2	0.0	0.0	18.7	58.8
2543	8.7	0.8	0.3	20.5	94.8
2544	12.5	1.1	0.4	20.4	91.6
2545	9.8	0.9	0.3	20.4	92.7
2546	21.0	1.9	0.6	20.4	89.8
2548	0.8	0.1	0.0	15.7	89.9
2549	1.8	0.2	0.1	20.4	96.7

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
2593	0.4	0.1	0.1	10.4	14.4
2599	1.1	0.2	0.2	10.7	16.8
2603	5.0	0.7	0.5	14.4	27.3
2604	0.2	0.0	0.0	14.3	31.1
2605	15.6	1.6	0.8	17.9	51.1
2606	2.0	0.3	0.2	14.1	26.6
2607	9.0	1.4	1.2	12.5	20.4
2608	32.1	3.4	1.8	17.6	47.9
2609	4.8	0.6	0.5	14.3	27.9
2610	11.6	1.5	1.2	14.0	26.6
2611	16.9	1.6	0.5	19.7	89.0
2612	12.5	1.7	1.3	13.8	25.6
2613	8.9	1.1	0.7	14.7	33.6
2614	2.0	0.2	0.1	17.8	54.2
2615	0.0	0.0	0.0	17.4	46.0
2616	4.0	0.4	0.2	19.1	69.0
2617	6.7	0.6	0.2	20.0	87.0
2618	22.4	2.0	0.6	20.6	96.3
2619	10.7	1.0	0.3	20.0	92.6
2620	1.8	0.2	0.1	19.6	77.3
2622	0.2	0.0	0.0	19.2	66.2
2623	0.0	0.0	0.0	19.1	63.9
2666	2.0	0.3	0.3	10.7	17.3
2669	0.0	0.0	0.0	NA	NA
2673	0.1	0.0	0.0	5.4	5.7
2674	1.4	0.2	0.2	10.7	16.0
2676	0.4	0.1	0.0	16.4	44.2
2677	2.6	0.3	0.2	14.4	29.7
2678	2.7	0.4	0.3	14.3	29.0
2679	28.5	3.6	2.6	14.9	29.7
2680	4.5	0.6	0.4	15.4	33.6
2681	5.5	0.7	0.6	13.8	24.8
2682	9.4	1.2	1.0	14.1	26.8
2684	7.0	0.8	0.4	16.9	47.0
2685	6.4	0.7	0.4	16.9	48.2
2686	5.3	0.6	0.4	16.3	40.5
2687	3.9	0.5	0.2	16.2	45.1

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
2688	0.4	0.0	0.0	17.2	47.8
2690	0.0	0.0	0.0	18.1	52.8
2691	5.5	0.6	0.2	17.0	63.0
2692	5.8	0.6	0.3	18.3	61.5
2743	0.0	0.0	0.0	NA	NA
2748	0.3	0.1	0.0	10.7	24.3
2749	3.2	0.4	0.3	14.4	32.1
2750	4.4	0.4	0.2	19.2	78.9
2751	7.5	0.8	0.4	17.7	56.7
2752	3.5	0.5	0.5	12.6	21.4
2753	8.7	1.1	0.8	14.9	30.6
2754	12.7	1.3	0.6	17.9	54.8
2755	19.5	2.4	1.7	15.1	31.3
2756	0.6	0.1	0.1	12.5	27.1
2757	11.1	1.4	1.0	15.2	32.0
2758	56.1	6.3	3.7	16.6	41.2
2759	11.7	1.2	0.5	18.5	70.0
2760	21.6	2.3	1.2	17.6	51.1
2761	4.0	0.4	0.2	18.0	62.1
2762	0.4	0.0	0.0	17.0	45.8
2765	0.7	0.1	0.0	17.8	49.2
2766	6.2	0.7	0.4	16.5	41.4
2767	5.5	0.8	0.6	13.7	26.0
2768	0.0	0.0	0.0	17.4	46.0
2816	1.3	0.2	0.2	10.9	17.3
2818	0.3	0.1	0.1	11.9	18.6
2821	0.0	0.0	0.0	NA	NA
2822	0.3	0.1	0.0	11.4	25.5
2824	5.3	0.6	0.4	15.2	36.1
2825	4.9	0.6	0.4	14.6	30.2
2826	4.2	0.6	0.5	13.3	25.3
2827	1.4	0.2	0.1	14.5	28.1
2829	7.0	1.0	0.8	13.2	23.6
2830	5.6	0.8	0.6	12.9	24.1
2831	10.9	1.4	1.0	14.6	31.3
2832	23.7	2.6	1.4	17.2	46.9
2833	24.7	2.6	1.3	17.9	52.9

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
2834	13.8	1.7	1.1	15.7	34.8
2835	0.0	0.0	0.0	17.2	44.3
2837	0.0	0.0	0.0	17.4	45.7
2839	0.0	0.0	0.0	18.0	51.8
2841	7.7	1.1	1.0	12.8	21.3
2842	0.5	0.1	0.0	16.5	38.7
2843	0.1	0.0	0.0	17.4	46.3
2897	0.0	0.0	0.0	NA	NA
2898	4.4	0.5	0.3	16.3	41.8
2899	0.4	0.1	0.0	15.2	33.4
2900	0.6	0.1	0.1	15.1	31.3
2902	10.3	1.5	1.2	13.2	24.3
2903	6.6	0.9	0.6	14.3	27.7
2904	3.3	0.4	0.3	15.5	35.5
2905	3.4	0.4	0.3	14.8	33.1
2906	49.9	5.6	3.3	16.7	41.2
2907	52.9	5.9	3.4	16.8	42.6
2908	27.4	3.7	1.7	13.9	44.3
2909	0.1	0.0	0.0	17.3	44.9
2910	0.9	0.1	0.0	17.9	52.2
2912	0.2	0.0	0.0	18.1	51.9
2913	0.8	0.1	0.0	18.0	57.6
2914	0.2	0.0	0.0	17.5	46.7
2915	0.7	0.1	0.0	17.3	47.7
2916	1.3	0.1	0.1	18.0	53.9
2962	0.2	0.0	0.0	11.2	18.2
2966	0.9	0.2	0.2	9.4	13.2
2971	0.0	0.0	0.0	12.0	18.8
2973	0.3	0.0	0.0	14.1	26.1
2974	7.8	1.0	0.8	14.1	26.3
2975	0.9	0.1	0.1	14.1	26.9
2976	2.8	0.4	0.3	13.8	26.7
2977	0.3	0.0	0.0	12.3	30.7
2978	0.1	0.0	0.0	15.6	33.5
2979	2.0	0.2	0.1	15.8	36.7
2980	15.5	1.7	0.9	17.4	46.3
2981	9.2	1.1	0.6	15.8	41.8

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
2982	13.8	1.5	0.8	16.9	46.8
2984	3.8	0.4	0.2	18.7	62.2
2985	6.1	0.6	0.2	20.1	86.4
2986	3.0	0.3	0.1	18.9	67.1
2987	2.7	0.3	0.1	17.6	64.8
2989	1.8	0.2	0.1	17.5	46.6
2990	0.1	0.0	0.0	17.9	50.1
2991	5.6	0.5	0.2	20.5	97.4
2992	0.0	0.0	0.0	18.0	50.9
3038	0.8	0.1	0.1	11.0	15.9
3040	5.0	1.3	1.3	7.1	10.6
3041	2.1	0.5	0.5	8.6	12.4
3043	14.6	2.5	2.6	10.8	15.6
3045	3.3	0.6	0.6	11.0	15.9
3046	1.6	0.3	0.3	11.1	16.3
3047	1.0	0.2	0.1	10.6	22.4
3048	1.1	0.1	0.1	14.1	26.0
3050	0.2	0.0	0.0	14.5	32.2
3051	0.0	0.0	0.0	15.4	32.5
3052	0.4	0.0	0.0	15.8	37.0
3053	3.9	0.4	0.2	16.5	51.2
3054	3.3	0.4	0.2	17.1	43.7
3055	4.1	0.4	0.2	18.7	63.2
3056	2.0	0.2	0.1	18.5	58.4
3057	7.6	0.8	0.4	18.4	56.7
3058	3.2	0.3	0.1	20.0	82.9
3059	9.1	0.9	0.4	19.3	70.1
3060	8.4	0.8	0.2	20.8	101.5
3061	0.5	0.1	0.0	18.1	56.8
3062	0.0	0.0	0.0	17.4	46.3
3067	0.1	0.0	0.0	16.7	40.6
3113	0.7	0.1	0.1	9.5	13.3
3117	0.9	0.2	0.2	9.6	13.7
3119	1.9	0.3	0.3	11.2	16.5
3120	0.7	0.1	0.1	10.5	15.1
3121	2.3	0.5	0.5	8.5	11.8
3123	0.0	0.0	0.0	14.2	26.8

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
3124	0.0	0.0	0.0	14.8	29.3
3125	0.0	0.0	0.0	15.3	31.6
3126	0.0	0.0	0.0	15.9	35.3
3127	0.1	0.0	0.0	16.6	39.6
3128	1.0	0.1	0.1	17.4	49.5
3129	2.9	0.3	0.1	19.1	66.5
3130	5.5	0.5	0.2	19.3	71.0
3131	5.4	0.5	0.2	18.5	61.1
3132	3.5	0.5	0.4	13.7	25.8
3133	2.3	0.2	0.1	18.5	62.3
3134	2.9	0.3	0.1	19.4	80.5
3136	0.7	0.1	0.0	17.0	44.9
3140	0.1	0.0	0.0	16.5	39.1
3188	0.2	0.0	0.1	7.6	8.9
3189	0.3	0.1	0.1	6.5	7.2
3191	1.4	0.5	0.7	5.0	5.4
3192	1.2	0.3	0.3	8.1	10.2
3196	1.9	0.3	0.2	13.2	22.5
3198	0.2	0.0	0.0	14.6	28.4
3199	2.5	0.4	0.3	13.1	22.3
3200	0.7	0.1	0.1	15.2	31.2
3201	4.2	0.5	0.3	16.4	38.2
3202	0.2	0.0	0.0	17.5	46.9
3203	2.0	0.2	0.1	18.4	59.4
3204	5.6	0.5	0.2	19.6	76.6
3205	13.7	1.7	1.3	14.7	29.7
3206	9.1	1.0	0.6	17.2	45.3
3207	9.3	1.0	0.5	17.7	49.3
3208	5.3	0.5	0.2	19.6	77.3
3213	0.0	0.0	0.0	16.1	36.6
3214	0.1	0.0	0.0	16.1	36.6
3215	0.1	0.0	0.0	16.1	36.4
3261	2.4	0.6	0.6	8.3	11.0
3262	1.1	0.4	0.4	5.6	7.1
3263	21.5	7.9	10.7	5.1	5.5
3264	25.5	10.1	14.4	4.7	4.9
3265	0.5	0.1	0.1	7.5	9.0

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
3266	4.0	0.9	1.0	8.5	10.7
3267	1.4	0.3	0.3	10.7	15.3
3268	3.1	0.5	0.5	11.3	17.7
3269	1.7	0.3	0.3	11.7	18.3
3270	0.2	0.0	0.0	13.9	29.3
3273	0.3	0.0	0.0	12.1	30.3
3274	1.5	0.2	0.1	15.4	32.5
3275	9.3	1.1	0.7	16.2	37.0
3276	3.2	0.4	0.2	16.9	44.7
3277	31.6	3.3	1.6	18.1	54.1
3278	8.0	0.8	0.3	19.2	70.9
3279	28.4	2.7	0.9	20.0	84.8
3280	7.3	1.1	0.9	12.6	21.5
3281	3.9	0.4	0.2	17.5	47.4
3282	17.9	1.7	0.6	20.1	87.6
3283	3.2	0.4	0.1	16.9	64.5
3284	2.1	0.4	0.4	10.4	16.0
3288	0.1	0.0	0.0	15.9	35.1
3289	0.2	0.0	0.0	15.9	35.1
3337	12.3	3.9	5.2	5.9	6.5
3338	1.8	0.5	0.7	6.7	7.7
3339	2.1	0.5	0.6	8.4	10.5
3340	0.6	0.1	0.1	9.8	13.2
3343	0.9	0.2	0.2	8.7	10.8
3344	0.9	0.1	0.1	12.8	23.5
3345	2.9	0.4	0.3	15.2	31.5
3346	0.5	0.1	0.0	16.6	45.6
3347	1.4	0.2	0.1	15.2	31.0
3348	6.6	0.7	0.4	17.1	43.5
3349	0.3	0.0	0.0	15.9	35.2
3350	9.7	1.1	0.7	16.3	38.3
3351	54.1	6.1	3.7	16.6	40.2
3352	50.0	4.8	1.9	19.5	72.4
3353	32.5	3.0	0.9	20.4	95.7
3354	22.9	2.8	1.0	15.5	64.7
3355	1.0	0.1	0.1	17.1	43.3
3356	6.4	0.7	0.2	17.5	83.8

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
3357	5.7	0.6	0.2	18.2	63.4
3358	1.0	0.1	0.1	13.1	40.7
3410	0.5	0.1	0.1	8.5	10.9
3412	0.0	0.0	0.0	NA	NA
3413	0.9	0.2	0.3	7.1	8.2
3414	1.2	0.2	0.3	8.9	11.4
3415	1.1	0.2	0.2	10.7	15.5
3416	2.7	0.4	0.4	11.7	18.1
3420	0.4	0.1	0.0	14.6	28.4
3421	0.3	0.0	0.0	14.7	28.9
3422	1.8	0.2	0.1	15.3	36.2
3423	1.4	0.2	0.1	14.9	29.9
3424	4.3	0.6	0.4	14.1	27.6
3425	27.7	3.8	3.1	13.5	24.5
3426	18.4	1.9	1.0	17.8	50.5
3427	11.0	1.1	0.4	19.6	79.9
3428	2.2	0.2	0.1	17.4	46.0
3429	8.9	1.4	1.3	11.9	18.8
3430	1.6	0.2	0.1	14.0	29.5
3431	2.6	0.3	0.2	15.4	34.8
3432	6.6	1.0	0.9	12.3	20.4
3433	2.9	0.5	0.4	10.7	17.8
3435	0.1	0.0	0.0	14.2	61.5
3485	0.6	0.1	0.2	7.5	9.2
3486	0.5	0.1	0.1	7.8	9.4
3487	2.9	0.7	0.8	7.5	9.9
3488	4.4	1.0	1.1	8.7	11.2
3489	0.7	0.1	0.1	10.1	15.2
3490	2.2	0.5	0.5	9.0	12.0
3492	0.1	0.0	0.0	13.2	22.8
3493	0.0	0.0	0.0	13.7	24.5
3494	0.0	0.0	0.0	14.1	26.2
3495	8.2	1.1	0.9	14.0	25.9
3496	11.0	1.6	1.5	12.6	20.7
3497	1.2	0.2	0.1	13.9	25.3
3498	25.4	3.5	2.8	13.7	24.6
3499	2.7	0.4	0.3	13.7	24.4

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
3500	0.7	0.1	0.1	15.5	33.2
3501	2.8	0.3	0.2	15.2	31.6
3502	5.0	0.7	0.5	14.1	26.2
3503	6.3	0.8	0.6	14.4	27.8
3504	17.7	2.4	1.9	13.8	25.1
3505	17.2	2.3	1.8	14.0	26.0
3506	21.5	2.9	2.4	13.7	24.8
3507	0.3	0.0	0.0	12.9	37.8
3508	1.2	0.1	0.1	16.6	42.1
3509	19.1	1.8	0.6	19.9	81.5
3559	7.1	1.7	2.0	7.8	9.5
3560	0.0	0.0	0.0	7.1	8.7
3561	1.6	0.5	0.6	6.4	7.1
3562	2.4	0.8	1.1	5.6	6.0
3563	1.0	0.2	0.3	8.9	11.3
3564	0.3	0.0	0.0	12.8	24.9
3568	2.6	0.3	0.3	14.0	26.2
3569	4.9	0.7	0.6	13.2	22.7
3570	3.7	0.5	0.5	12.9	21.9
3571	0.7	0.1	0.1	11.9	24.2
3572	24.0	3.3	2.8	13.4	23.4
3573	14.1	2.1	1.8	12.7	21.1
3574	3.0	0.4	0.3	13.9	25.2
3575	1.0	0.1	0.1	14.3	27.2
3576	5.7	0.8	0.6	14.2	26.5
3577	6.3	0.8	0.7	14.1	26.4
3578	2.8	0.4	0.3	14.1	26.2
3579	8.7	1.2	0.9	14.0	25.8
3580	14.7	2.0	1.6	13.8	25.6
3581	1.6	0.2	0.1	16.8	42.2
3582	2.8	0.3	0.2	17.3	48.4
3583	4.9	0.6	0.3	15.7	44.4
3633	0.6	0.2	0.2	7.4	8.6
3634	4.4	1.3	1.8	6.0	6.5
3635	4.2	1.1	1.5	6.9	7.8
3636	0.6	0.1	0.1	9.0	11.7
3637	0.2	0.0	0.0	11.6	19.3

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
3640	0.1	0.0	0.0	11.9	18.4
3641	2.7	0.6	0.8	8.1	10.0
3642	0.1	0.0	0.0	12.9	21.8
3644	0.0	0.0	0.0	12.9	21.5
3645	8.9	1.7	1.8	9.9	13.3
3646	5.4	0.8	0.6	12.9	22.9
3647	11.0	1.7	1.5	12.2	19.6
3648	21.0	3.1	2.8	12.6	20.7
3649	10.2	1.4	1.2	13.2	22.8
3650	10.8	1.5	1.3	13.2	22.8
3651	2.0	0.3	0.2	13.7	24.5
3652	3.5	0.5	0.4	13.6	24.8
3653	6.8	0.9	0.8	13.7	24.9
3654	3.2	0.5	0.4	12.8	25.0
3655	4.2	0.5	0.4	14.6	28.6
3656	3.5	0.4	0.3	16.1	37.3
3657	3.5	0.4	0.2	17.4	47.9
3658	0.1	0.0	0.0	15.7	34.2
3659	0.0	0.0	0.0	15.3	31.6
3707	0.3	0.1	0.1	9.1	11.6
3708	0.0	0.0	0.0	7.5	8.8
3709	0.4	0.1	0.1	7.3	10.6
3710	0.8	0.1	0.2	10.1	13.8
3711	1.6	0.2	0.1	18.3	66.6
3712	3.6	0.4	0.2	18.1	60.5
3713	0.6	0.1	0.1	11.4	17.4
3717	0.6	0.1	0.1	12.8	21.4
3718	4.1	0.7	0.6	11.7	18.0
3719	12.1	1.9	1.7	12.1	19.2
3720	43.4	6.7	6.1	12.1	19.6
3721	18.9	2.8	2.5	12.6	21.1
3722	3.8	0.6	0.5	12.8	21.4
3723	18.2	2.9	2.7	11.8	18.4
3724	21.0	3.2	2.9	12.4	20.2
3725	19.2	2.8	2.5	12.6	21.1
3726	5.3	0.8	0.7	12.3	19.6
3727	1.3	0.2	0.2	13.3	23.0

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
3728	0.2	0.0	0.0	13.7	24.6
3729	1.4	0.2	0.1	15.3	32.5
3730	1.1	0.1	0.1	14.4	29.4
3731	6.8	0.8	0.5	15.2	35.1
3780	40.1	3.1	0.1	24.6	1160.8
3783	4.3	1.2	1.5	6.9	7.8
3786	0.6	0.1	0.1	13.4	30.0
3787	0.7	0.1	0.1	11.3	17.4
3789	0.1	0.0	0.0	12.8	21.3
3791	1.2	0.2	0.2	12.8	21.5
3792	1.9	0.3	0.3	11.8	18.3
3793	7.5	1.3	1.3	10.8	15.7
3794	23.6	3.5	3.1	12.6	20.8
3795	60.9	9.3	8.4	12.3	19.8
3796	4.0	0.6	0.5	12.6	21.7
3797	7.2	1.1	0.9	12.5	20.8
3798	21.3	3.2	3.0	12.3	19.7
3799	25.8	4.1	3.9	11.8	18.3
3800	35.3	5.4	4.9	12.3	19.7
3801	36.2	5.6	5.3	12.0	18.9
3802	5.4	0.8	0.6	13.2	22.8
3803	4.7	0.7	0.5	12.7	24.4
3804	0.3	0.0	0.0	13.8	25.9
3805	1.0	0.1	0.1	13.8	27.6
3855	0.1	0.0	0.0	11.0	16.2
3864	1.2	0.2	0.2	12.9	22.1
3865	17.7	2.1	1.4	15.7	34.4
3866	12.7	2.0	1.8	12.0	19.0
3867	0.9	0.1	0.1	12.0	18.8
3868	5.2	0.8	0.7	12.7	20.9
3869	10.1	1.5	1.3	12.6	20.8
3870	3.1	0.5	0.4	12.5	20.7
3871	3.5	0.5	0.5	12.7	20.9
3872	33.7	5.2	4.9	12.0	19.0
3873	39.9	6.2	5.7	12.1	19.1
3874	88.8	13.3	12.0	12.5	20.3
3875	49.8	7.8	7.3	12.0	18.6

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
3876	39.1	4.4	2.7	16.5	39.6
3877	17.4	2.5	2.2	12.9	21.9
3878	0.6	0.1	0.1	13.7	24.4
3936	1.3	0.2	0.2	11.8	21.2
3938	0.3	0.0	0.0	12.9	21.6
3939	1.4	0.2	0.2	13.0	21.8
3940	2.2	0.3	0.3	12.5	21.2
3941	1.0	0.1	0.1	12.7	21.0
3944	2.7	0.4	0.4	12.5	20.2
3945	13.9	2.1	1.9	12.5	20.6
3946	17.8	2.6	2.2	12.9	22.3
3947	16.2	2.4	2.2	12.5	20.2
3948	53.5	7.9	7.1	12.6	20.8
3949	20.8	3.0	2.7	12.9	21.5
3950	6.5	1.0	0.9	12.5	20.4
4000	0.1	0.0	0.0	12.0	18.9
4004	5.9	0.9	0.9	11.7	18.1
4008	0.1	0.0	0.0	12.4	20.0
4012	0.5	0.1	0.1	12.7	21.0
4014	0.3	0.0	0.0	12.7	20.9
4018	1.7	0.2	0.2	12.8	21.4
4019	0.1	0.0	0.0	12.9	23.0
4020	3.9	0.6	0.5	13.0	22.0
4021	3.7	0.5	0.5	12.9	22.1
4022	8.3	1.2	1.1	12.7	21.0
4023	2.1	0.3	0.3	12.9	21.5
4024	0.3	0.0	0.0	13.0	22.1
4081	0.6	0.1	0.1	12.4	20.0
4085	0.2	0.0	0.0	12.5	20.3
4087	2.0	0.3	0.3	12.5	20.1
4088	1.8	0.3	0.2	12.4	20.0
4092	6.9	1.0	0.9	12.7	20.9
4093	1.0	0.1	0.1	13.4	23.5
4094	7.7	1.1	0.9	13.6	24.7
4095	19.7	2.5	1.8	14.8	29.5
4096	6.6	1.4	1.6	8.8	11.1
4158	0.5	0.1	0.1	12.3	19.7

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
4159	5.1	0.9	0.7	10.5	20.1
4160	0.6	0.1	0.1	12.0	19.3
4162	0.7	0.1	0.1	11.8	18.8
4163	15.5	2.5	2.3	11.9	18.4
4164	2.9	0.5	0.4	11.8	18.1
4165	1.9	0.3	0.3	11.7	18.1
4166	2.9	0.5	0.4	11.9	18.8
4167	8.6	1.4	1.3	11.5	17.8
4168	11.6	1.7	1.5	12.7	21.4
4169	9.1	1.3	1.2	12.7	21.1
4228	0.2	0.0	0.0	12.4	19.9
4229	0.0	0.0	0.0	12.3	19.8
4235	1.1	0.2	0.2	11.6	17.6
4236	3.7	0.6	0.6	11.3	17.1
4237	3.3	0.6	0.6	11.2	16.4
4238	2.6	0.5	0.5	10.4	15.5
4240	0.3	0.1	0.1	10.8	15.5
4241	12.9	2.1	2.1	11.3	16.8
4242	2.9	0.5	0.4	11.6	17.6
4302	0.7	0.1	0.1	12.2	19.5
4306	6.6	1.0	1.0	11.9	18.4
4307	1.9	0.3	0.3	11.5	17.6
4311	5.4	1.0	1.0	10.3	14.3
4313	1.7	0.4	0.5	7.5	9.0
4314	0.8	0.2	0.2	8.7	11.0
4315	10.1	2.1	2.4	9.0	11.6
4376	0.7	0.1	0.1	12.1	19.2
4382	0.0	0.0	0.0	NA	NA
4383	1.3	0.2	0.3	10.0	13.5
4384	0.6	0.1	0.1	9.5	14.6
4385	4.6	0.9	1.0	9.3	12.5
4386	0.1	0.0	0.0	7.2	11.4
4387	0.9	0.2	0.3	8.0	9.6
4388	2.6	0.7	1.0	6.4	7.1
4389	13.2	4.6	6.5	5.3	5.5
4442	1.2	0.2	0.2	12.3	20.0
4443	0.0	0.0	0.0	12.3	19.6

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
4458	10.7	3.0	3.8	6.7	7.7
4459	2.6	0.6	0.7	8.2	10.2
4460	1.9	0.4	0.5	8.0	9.7
4461	4.7	1.2	1.5	7.3	8.5
4462	8.5	2.8	3.9	5.7	6.0
4463	2.5	0.7	0.8	7.0	7.9
4516	0.0	0.0	0.0	NA	NA
4522	0.7	0.1	0.1	11.3	16.9
4527	2.7	0.3	0.2	15.6	48.8
4528	0.0	0.0	0.0	12.8	21.3
4529	0.0	0.0	0.0	NA	NA
4531	0.4	0.1	0.1	7.9	9.6
4532	0.1	0.0	0.1	6.1	6.8
4533	1.6	0.4	0.5	6.8	8.2
4534	5.1	1.8	2.6	5.2	5.4
4535	3.4	0.9	1.2	7.0	8.1
4536	4.5	1.3	1.7	6.3	7.2
4537	0.0	0.0	0.0	7.1	8.1
4543	0.7	0.1	0.1	12.3	19.7
4590	0.3	0.1	0.0	12.4	19.9
4594	1.1	0.3	0.3	7.2	9.0
4595	0.0	0.0	0.0	7.8	9.3
4597	0.5	0.1	0.1	11.2	18.1
4599	0.0	0.0	0.0	NA	NA
4600	1.1	0.2	0.2	11.7	18.5
4605	1.2	0.3	0.4	6.8	7.8
4606	4.4	1.3	1.6	6.6	7.6
4607	25.7	9.3	12.5	5.2	5.7
4608	4.5	1.6	2.2	5.4	5.7
4609	9.4	1.9	2.2	9.1	11.7
4610	1.8	0.5	0.6	6.7	7.5
4672	0.3	0.1	0.1	11.8	18.2
4674	1.4	0.2	0.2	11.6	17.5
4679	1.1	0.4	0.5	5.4	5.7
4680	4.5	1.7	2.3	4.9	5.5
4682	13.0	4.7	6.6	5.2	5.4
4683	8.7	3.2	4.5	5.1	5.3

Table E-2 (continued).

Grid Cell ID	2009 Methane emissions (kg/h)	2009 Ethane emissions (kg/h)	2009 Propane emissions (kg/h)	Methane to Ethane ratio	Methane to Propane ratio
4684	1.9	0.6	0.7	6.5	7.2
4753	1.1	0.3	0.3	7.8	9.3
4756	1.0	0.3	0.4	6.3	7.0
4757	1.2	0.4	0.5	6.0	6.5
4758	0.1	0.0	0.0	6.6	7.4

For each grid cell, Table S3-3 shows EF factor, activity factor and estimated 2013 methane emissions.

Table E-3. Estimated methane 2013 emissions, based on emission factor (methane emissions 2009 / natural gas production 2009) and activity factor (natural gas production 2013) per grid cell.

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
1066	NA	NA	NA
1139	0.0002	300.8	0.1
1285	0.0006	878.5	1.1
1287	0.0001	229.2	0.0
1288	0.0031	1368.4	9.3
1359	0.0004	160.9	0.1
1360	0.0087	781.6	14.8
1361	0.0017	325.8	1.2
1364	0.0000	339.9	0.0
1416	0.0792	0.0	0.0
1428	0.0954	113.1	23.7
1433	0.0024	15.9	0.1
1434	0.0011	606.9	1.5
1435	0.0006	204.1	0.3
1436	0.0010	409.3	0.9
1437	0.0002	286.4	0.1
1493	0.0334	0.0	0.0
1494	0.4181	0.0	0.0
1496	0.0495	42.4	4.6
1498	NA	NA	NA
1504	0.0197	101.7	4.4
1505	0.0100	253.8	5.6
1506	0.0013	73.8	0.2
1507	0.0033	191.7	1.4
1508	0.0022	353.9	1.7
1509	0.0005	649.5	0.7
1510	0.0045	644.6	6.4
1511	0.0031	467.0	3.2
1557	NA	0.0	NA
1558	NA	0.0	NA
1566	0.0018	0.0	0.0
1568	NA	274.7	NA

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
1569	0.0289	0.0	0.0
1574	NA	NA	NA
1575	NA	2246.4	NA
1580	0.0027	902.2	5.4
1581	0.0011	242.7	0.6
1582	0.0045	298.0	3.0
1583	0.0020	484.7	2.1
1584	0.0014	538.1	1.7
1639	0.0040	0.0	0.0
1640	0.0033	0.0	0.0
1642	NA	54.3	NA
1646	0.0011	0.0	0.0
1647	0.0405	947.4	84.1
1648	0.0624	0.0	0.0
1649	0.0127	1346.1	37.4
1650	0.1984	3585.8	1559.5
1651	0.0015	10735.0	35.3
1652	0.0145	3928.6	124.9
1653	0.0062	560.3	7.6
1654	0.0045	4428.8	43.4
1655	0.0023	556.6	2.8
1656	0.0057	70.7	0.9
1657	0.0009	1532.8	3.0
1658	0.0044	1251.8	12.1
1659	0.7271	345.5	550.5
1717	0.0151	35.2	1.2
1718	0.0275	0.0	0.0
1722	NA	1279.3	NA
1723	0.0069	635.3	9.6
1724	0.0015	1410.9	4.6
1725	0.0001	3156.3	0.7
1726	0.0008	1425.9	2.4
1727	0.0080	3151.1	55.4
1728	0.0156	6783.5	231.3
1729	0.0010	1014.2	2.1
1730	0.0000	449.6	0.0
1731	0.0000	912.2	0.1
1732	0.0003	1488.2	0.8

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
1786	NA	56.7	NA
1787	0.1256	23.9	6.6
1789	0.0024	0.0	0.0
1790	0.0091	0.0	0.0
1794	0.0037	1379.9	11.2
1795	0.0034	527.7	3.9
1796	0.0028	2030.4	12.3
1797	0.0022	1816.7	8.8
1798	0.0004	951.5	0.8
1799	0.0029	2475.7	15.8
1800	0.0002	1728.5	0.8
1801	0.0002	2273.4	0.8
1802	0.0005	3947.8	4.5
1803	0.0027	2278.6	13.3
1804	0.0001	1113.3	0.2
1805	0.0001	738.5	0.2
1806	0.0013	3243.0	9.6
1809	0.0003	679.0	0.5
1859	0.0028	396.4	2.4
1862	NA	127.0	NA
1863	0.0222	0.0	0.0
1864	0.0236	0.0	0.0
1865	0.0092	0.0	0.0
1868	0.0224	698.6	34.3
1869	NA	NA	NA
1870	0.0011	2624.9	6.3
1871	0.0005	2309.4	2.5
1872	0.0003	3133.7	2.0
1873	0.0004	5702.6	5.3
1874	0.0001	2849.9	0.7
1875	0.0001	4470.7	1.2
1876	0.0001	7229.7	1.1
1877	0.0002	2531.4	1.3
1878	0.0001	7251.8	1.6
1879	0.0008	1169.9	1.9
1880	0.0017	4254.5	15.5
1881	0.0043	2847.3	27.0
1883	0.0004	506.9	0.5

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
1932	0.0012	39.8	0.1
1933	0.0025	518.5	2.8
1938	0.0202	53.1	2.4
1941	0.0049	0.0	0.0
1942	0.0041	2282.6	20.3
1943	0.0064	1986.0	27.6
1944	0.0394	2138.3	184.9
1945	0.0015	3749.2	12.3
1946	0.0003	1455.1	1.0
1947	0.0003	3456.8	2.0
1948	0.0000	1913.3	0.1
1949	0.0008	2603.1	4.3
1950	0.0001	2138.4	0.3
1951	0.0002	3763.8	1.7
1952	0.0002	8880.6	4.6
1953	0.0000	6404.7	0.6
1954	0.0220	9203.5	444.8
1955	0.0084	8981.7	164.8
1956	0.0053	5407.3	62.6
2005	0.0054	0.0	0.0
2006	0.0019	0.0	0.0
2007	0.0047	103.2	1.1
2008	0.0096	130.9	2.8
2009	NA	255.0	NA
2011	0.0052	0.0	0.0
2012	0.0054	0.0	0.0
2013	0.0460	392.3	39.6
2014	0.0129	0.0	0.0
2015	0.0044	460.0	4.5
2016	0.0067	655.6	9.6
2017	0.0033	2799.8	20.5
2018	0.0050	2478.9	27.1
2019	0.0028	3815.6	23.4
2020	0.0002	2688.6	1.2
2021	0.0001	969.8	0.1
2022	0.0002	3783.0	1.3
2023	0.0000	906.6	0.1
2024	0.0006	4984.4	6.3

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
2025	0.0058	3953.7	49.8
2026	0.0001	6456.9	0.8
2027	0.0001	7608.2	0.8
2028	0.0036	11717.2	92.2
2029	0.0085	8808.0	164.3
2030	0.1769	3960.4	1535.7
2072	0.0483	84.3	8.9
2073	0.0052	76.7	0.9
2080	0.0019	45.3	0.2
2081	0.0019	58.1	0.2
2082	0.0020	126.7	0.6
2083	0.0035	151.5	1.2
2084	0.0020	0.0	0.0
2086	0.0439	0.0	0.0
2087	0.0085	1022.2	19.1
2088	0.0112	481.0	11.9
2089	0.0033	715.4	5.2
2090	0.0030	469.5	3.0
2091	0.0033	5356.6	39.1
2092	0.0014	2748.3	8.3
2093	0.0004	4112.8	3.4
2094	0.0007	3991.5	6.3
2095	0.0000	682.5	0.1
2096	0.0000	3044.7	0.1
2097	0.0000	4209.1	0.3
2098	0.0007	3191.7	5.0
2099	0.0002	7569.1	4.0
2100	0.0006	10756.4	13.0
2101	0.0003	13484.9	8.7
2102	0.0007	9044.4	13.0
2103	0.0022	2307.1	10.9
2104	0.0006	3365.9	4.1
2147	0.0548	81.2	9.8
2149	0.0000	4.8	0.0
2154	0.0019	23.5	0.1
2155	0.0009	0.0	0.0
2156	0.0021	0.0	0.0
2158	0.0139	1.1	0.0

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
2159	NA	0.0	NA
2160	0.0056	0.0	0.0
2162	NA	0.0	NA
2163	0.0054	809.7	9.5
2165	0.0027	1193.5	7.1
2166	0.0019	100.4	0.4
2167	0.0009	1484.3	2.9
2168	0.0003	2879.2	1.8
2169	0.0001	325.2	0.0
2170	0.0000	166.1	0.0
2171	0.0008	6367.8	11.0
2172	0.0003	4075.0	2.9
2173	0.0019	6470.0	27.3
2174	0.0003	27955.3	21.4
2175	0.0020	9713.2	43.4
2176	0.0044	10613.9	102.1
2177	0.0017	8609.9	32.2
2178	0.0181	4355.4	172.6
2179	0.0230	6381.3	321.8
2225	0.4873	0.0	0.0
2227	NA	0.0	NA
2228	0.0183	0.0	0.0
2229	0.0012	11.3	0.0
2230	0.0026	0.0	0.0
2233	0.0060	0.0	0.0
2234	0.0026	0.0	0.0
2235	NA	0.0	NA
2236	0.0052	0.0	0.0
2237	0.0012	106.0	0.3
2238	0.0028	227.8	1.4
2239	0.0048	1486.9	15.5
2240	0.0023	1575.9	8.0
2241	0.0010	3168.0	6.9
2242	0.0018	4294.4	17.4
2243	0.0012	1183.7	3.0
2244	0.0008	6007.8	10.4
2245	0.0224	4021.6	197.0
2246	0.0008	4359.3	7.3

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
2247	0.0003	18740.9	11.5
2248	0.0018	10386.7	41.1
2249	0.0009	7243.5	14.0
2250	0.0008	6549.5	11.2
2251	0.0109	8177.6	195.9
2252	0.0014	3025.2	9.6
2294	0.0089	0.0	0.0
2296	0.0823	10.3	1.9
2297	NA	NA	NA
2299	0.4413	0.0	0.0
2301	0.0067	0.0	0.0
2302	0.4935	0.0	0.0
2303	0.0031	13.0	0.1
2304	0.1069	0.0	0.0
2305	NA	0.0	NA
2307	0.0521	0.0	0.0
2308	0.0011	66.8	0.2
2309	0.0011	23.4	0.1
2310	0.0053	1158.7	13.5
2311	0.0006	715.1	1.0
2312	0.0014	299.2	0.9
2313	0.0025	440.6	2.5
2314	0.0021	282.4	1.3
2315	0.0008	1002.1	1.6
2316	0.0023	7510.2	37.2
2317	0.0006	9125.4	11.0
2318	0.0004	9807.7	7.9
2319	0.0001	963.8	0.2
2320	0.0009	6024.2	11.9
2321	0.0018	2814.8	11.0
2322	0.0030	2053.2	13.4
2323	0.0009	3930.0	7.8
2324	0.0024	4664.1	24.9
2325	0.0025	3891.9	21.0
2326	0.0211	14056.5	648.7
2328	0.0021	4873.1	22.8
2368	0.0040	0.0	0.0
2374	0.0001	0.0	0.0

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
2379	0.0053	0.0	0.0
2380	0.0040	0.0	0.0
2381	0.0065	0.0	0.0
2382	0.0072	45.4	0.7
2383	0.0042	537.1	4.9
2384	0.0077	491.4	8.3
2385	0.0042	683.3	6.3
2386	0.0003	1276.1	0.8
2387	0.0025	455.6	2.5
2388	0.0011	391.2	1.0
2389	0.0032	2394.4	16.6
2390	0.0017	4257.7	16.2
2391	0.0010	3887.8	8.5
2392	0.0026	15217.1	86.1
2393	0.0002	1479.8	0.7
2394	0.0003	3048.3	1.7
2395	0.0007	1986.3	3.0
2396	0.0004	13119.0	10.3
2397	0.0003	4231.2	2.8
2398	0.0015	9100.1	30.8
2399	0.0061	20822.6	278.7
2400	0.0044	11556.7	112.5
2401	0.0010	5500.7	12.4
2402	0.0000	2368.0	0.1
2448	NA	0.0	NA
2449	NA	0.0	NA
2454	0.0000	0.0	0.0
2456	0.0205	0.0	0.0
2458	0.0013	840.8	2.4
2459	0.0024	3526.4	18.3
2460	0.0019	202.6	0.8
2461	0.0045	564.3	5.5
2462	0.0007	67.7	0.1
2463	0.0018	2281.4	8.9
2465	0.0004	1465.6	1.2
2466	0.0001	6230.2	1.6
2467	0.0000	10074.6	0.8
2468	0.0000	11040.7	0.2

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
2469	0.0002	5769.1	2.3
2470	0.0005	3252.7	3.2
2471	0.0003	5371.1	3.8
2472	0.0004	8277.9	7.7
2473	0.0044	8562.8	83.4
2474	0.0014	7375.0	23.1
2475	0.0001	11447.6	1.6
2518	0.0015	0.0	0.0
2519	0.0030	0.0	0.0
2523	0.0018	0.0	0.0
2528	0.0024	0.0	0.0
2529	0.0035	172.9	1.3
2530	0.1009	62.1	13.7
2531	0.0013	92.7	0.3
2532	0.0083	175.9	3.2
2533	0.0041	1192.0	10.6
2534	0.0612	1996.6	267.8
2535	0.0018	1807.3	7.0
2536	0.0028	68.1	0.4
2537	0.0019	4417.0	18.8
2538	0.0007	4278.0	6.4
2539	0.0009	1728.6	3.2
2540	0.0003	2645.9	1.7
2541	0.0003	2398.5	1.4
2542	0.0000	6492.0	0.2
2543	0.0003	4700.0	2.7
2544	0.0007	1867.6	2.7
2545	0.0003	3416.3	2.3
2546	0.0005	11781.1	14.1
2548	0.0001	3880.7	0.8
2549	0.0002	5678.8	2.2
2593	0.0281	0.0	0.0
2599	0.0038	0.0	0.0
2603	0.0118	105.4	2.7
2604	NA	NA	NA
2605	0.0069	148.5	2.2
2606	0.0019	382.2	1.6
2607	0.0098	140.4	3.0

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
2608	0.0112	0.0	0.0
2609	0.0044	136.2	1.3
2610	0.0043	350.8	3.3
2611	0.0020	2078.0	9.3
2612	0.0008	6389.2	11.3
2613	0.0007	4352.1	6.6
2614	0.0003	2364.1	1.4
2615	0.0000	2391.5	0.0
2616	0.0014	13083.2	38.7
2617	0.0003	6755.4	4.9
2618	0.0009	5039.0	10.2
2619	0.0004	9985.5	7.8
2620	0.0001	1454.7	0.4
2622	0.0000	3618.8	0.1
2623	NA	5260.0	NA
2666	NA	0.0	NA
2669	NA	0.0	NA
2673	NA	0.0	NA
2674	0.0047	0.0	0.0
2676	0.0123	0.0	0.0
2677	0.0196	0.0	0.0
2678	0.0048	107.3	1.1
2679	0.0074	0.0	0.0
2680	0.0044	51.3	0.5
2681	0.0045	61.4	0.6
2682	0.0056	3593.2	43.7
2684	0.0016	208.2	0.7
2685	0.0015	278.2	0.9
2686	0.0022	939.4	4.5
2687	0.0005	5032.5	5.5
2688	0.0001	5338.1	1.6
2690	0.0000	4456.4	0.1
2691	0.0003	4241.4	2.9
2692	0.0010	7433.9	15.9
2743	NA	0.0	NA
2748	0.0069	0.0	0.0
2749	0.0100	0.0	0.0
2750	0.0061	0.0	0.0

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
2751	0.0029	10.9	0.1
2752	0.0064	0.0	0.0
2753	0.0086	86.0	1.6
2754	0.0051	170.5	1.9
2755	0.0041	0.0	0.0
2756	0.0007	83.3	0.1
2757	0.0028	20.2	0.1
2758	0.0049	0.0	0.0
2759	0.0018	2166.6	8.4
2760	0.0034	3762.1	28.1
2761	0.0008	739.5	1.2
2762	0.0007	1602.7	2.5
2765	0.0001	12140.2	2.4
2766	0.0017	8721.1	33.0
2767	0.0003	8946.0	6.1
2768	0.0000	4379.4	0.2
2816	0.0081	0.0	0.0
2818	NA	0.0	NA
2821	NA	46.0	NA
2822	0.0036	99.7	0.8
2824	0.0026	0.0	0.0
2825	0.0035	159.3	1.2
2826	0.0039	48.7	0.4
2827	0.0043	0.4	0.0
2829	0.0044	0.0	0.0
2830	0.0081	82.1	1.5
2831	0.0057	1526.4	19.1
2832	0.0069	373.0	5.6
2833	0.0034	1122.1	8.3
2834	0.0027	2012.5	11.9
2835	0.0000	1973.9	0.1
2837	NA	5808.7	NA
2839	0.0000	5.3	0.0
2841	0.0006	354.1	0.5
2842	0.0006	9411.3	12.0
2843	0.0000	2570.0	0.0
2897	NA	21.6	NA
2898	0.0120	0.0	0.0

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
2899	0.0012	156.3	0.4
2900	0.0006	799.8	1.1
2902	0.0044	62.3	0.6
2903	0.0016	156.7	0.5
2904	0.0047	0.0	0.0
2905	0.0031	10.2	0.1
2906	0.0027	39.6	0.2
2907	0.0038	113.9	0.9
2908	0.0011	338.5	0.8
2909	0.0000	2752.0	0.2
2910	0.0003	4269.4	2.8
2912	0.0000	5881.9	0.3
2913	0.0001	2027.0	0.4
2914	0.0000	260.1	0.0
2915	0.0001	0.0	0.0
2916	0.0003	1707.7	1.1
2962	0.0196	0.0	0.0
2966	0.0081	140.8	2.5
2971	0.0003	326.4	0.2
2973	0.0030	0.0	0.0
2974	0.0094	42.7	0.9
2975	0.0032	0.0	0.0
2976	0.0154	52.5	1.8
2977	0.0007	0.0	0.0
2978	0.0004	153.3	0.1
2979	0.0011	210.4	0.5
2980	0.0044	30.5	0.3
2981	0.0025	627.1	3.4
2982	0.0014	562.0	1.7
2984	0.0004	9109.6	8.3
2985	0.0004	9458.9	8.6
2986	0.0004	7916.4	6.9
2987	0.0009	7337.1	14.8
2989	0.0006	11.2	0.0
2990	0.0000	121.1	0.0
2991	0.0003	0.0	0.0
2992	0.0017	0.0	0.0
3038	0.0060	0.0	0.0

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
3040	0.0121	0.0	0.0
3041	0.0047	30.8	0.3
3043	0.7531	1237.8	2043.2
3045	NA	177.2	NA
3046	0.0060	56.7	0.7
3047	0.0090	191.5	3.8
3048	0.0392	88.5	7.6
3050	0.0178	54.6	2.1
3051	0.0002	119.6	0.0
3052	0.0001	846.0	0.2
3053	0.0022	157.1	0.8
3054	0.0018	938.9	3.8
3055	0.0005	3823.5	4.4
3056	0.0020	3186.2	14.1
3057	0.0006	7047.0	9.0
3058	0.0027	8680.5	52.2
3059	0.0012	12467.1	33.0
3060	0.0027	7954.4	46.6
3061	0.0007	10338.9	15.4
3062	0.0000	14513.1	0.6
3067	0.0001	0.0	0.0
3113	0.0064	0.0	0.0
3117	0.0077	525.7	8.8
3119	NA	94.0	NA
3120	0.0025	0.0	0.0
3121	0.0025	2.6	0.0
3123	0.0003	0.0	0.0
3124	0.0001	0.0	0.0
3125	NA	246.0	NA
3126	0.0001	304.3	0.1
3127	0.0002	152.4	0.1
3128	0.0005	704.2	0.8
3129	0.0005	3491.7	4.1
3130	0.0009	1717.9	3.6
3131	0.0002	4692.2	2.1
3132	0.0003	3202.2	2.4
3133	0.0004	11896.9	10.3
3134	0.0045	9962.0	99.1

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
3136	0.0004	7622.1	7.2
3140	0.0000	0.0	0.0
3188	0.0145	0.0	0.0
3189	0.0270	0.0	0.0
3191	0.0022	0.0	0.0
3192	0.0069	14.8	0.2
3196	0.0076	93.2	1.6
3198	0.0009	0.0	0.0
3199	0.0168	390.2	14.3
3200	0.0018	326.2	1.3
3201	0.0087	81.1	1.6
3202	0.0001	55.9	0.0
3203	0.0004	641.0	0.5
3204	0.0003	2903.3	1.8
3205	0.0004	5358.9	4.5
3206	0.0002	1681.2	0.9
3207	0.0004	2691.7	2.6
3208	0.0020	7661.8	34.4
3213	0.0000	58.1	0.0
3214	0.0000	478.4	0.0
3215	0.0000	4775.3	0.2
3261	0.0102	0.0	0.0
3262	0.0040	0.0	0.0
3263	0.0018	0.0	0.0
3264	0.0114	43.7	1.1
3265	0.0040	17.0	0.1
3266	0.0047	200.7	2.1
3267	0.0102	139.5	3.1
3268	0.0039	284.3	2.5
3269	0.0029	195.3	1.3
3270	0.0007	297.3	0.5
3273	0.0024	210.5	1.1
3274	0.0107	184.0	4.3
3275	0.0269	1183.7	69.7
3276	0.0010	1763.7	4.0
3277	0.0024	1863.1	9.9
3278	0.0003	7540.9	4.6
3279	0.0007	8897.8	13.8

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
3280	0.0002	8355.0	3.9
3281	0.0001	11859.2	2.7
3282	0.0007	7826.1	12.2
3283	0.0008	5267.1	8.7
3284	0.0009	3410.0	7.0
3288	0.0000	1545.6	0.0
3289	0.0000	487.9	0.0
3337	0.0562	188.1	23.2
3338	NA	44.5	NA
3339	0.0092	0.0	0.0
3340	0.0051	137.4	1.5
3343	0.0036	0.0	0.0
3344	0.1379	10.6	3.2
3345	0.0093	0.0	0.0
3346	0.0210	0.0	0.0
3347	0.0022	148.6	0.7
3348	0.0220	471.2	22.7
3349	0.0007	668.3	1.0
3350	0.0044	949.0	9.1
3351	0.0049	1312.5	14.1
3352	0.0028	7787.8	47.0
3353	0.0011	7452.3	18.1
3354	0.0009	8293.6	16.0
3355	0.0000	8180.1	0.6
3356	0.0002	9300.6	4.4
3357	0.0005	8640.2	10.1
3358	0.0007	10648.4	16.3
3410	NA	25.2	NA
3412	NA	0.0	NA
3413	0.0010	26.7	0.1
3414	0.0022	205.5	1.0
3415	0.0115	218.7	5.5
3416	0.0045	44.0	0.4
3420	0.0014	88.9	0.3
3421	0.0008	272.6	0.5
3422	0.0062	911.4	12.4
3423	0.0135	445.3	13.2
3424	0.0041	849.1	7.6

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
3425	0.0040	3064.2	26.7
3426	0.0046	2194.7	22.3
3427	0.0015	5007.7	16.8
3428	0.0001	10343.6	1.8
3429	0.0003	5301.5	3.7
3430	0.0001	6859.9	1.1
3431	0.0003	8700.8	6.4
3432	0.0016	9597.9	32.6
3433	0.0011	9841.2	23.5
3435	0.0002	943.2	0.5
3485	0.0071	137.2	2.1
3486	0.0037	53.3	0.4
3487	0.0014	190.8	0.6
3488	0.0033	186.0	1.4
3489	0.0079	306.7	5.3
3490	0.0026	68.3	0.4
3492	0.0010	11.7	0.0
3493	0.0001	0.0	0.0
3494	0.0002	0.0	0.0
3495	0.0056	0.0	0.0
3496	0.0062	447.6	6.1
3497	0.0576	1663.5	210.0
3498	0.0051	1627.4	18.3
3499	0.0002	1083.8	0.4
3500	0.0000	3173.0	0.3
3501	0.0001	4120.8	1.2
3502	0.0004	2499.6	2.2
3503	0.0002	1604.0	0.8
3504	0.0016	2918.5	10.5
3505	0.0014	2731.0	8.1
3506	0.0041	4351.0	38.7
3507	0.0003	7266.2	5.4
3508	0.0005	6861.3	8.1
3509	0.0031	5871.2	40.0
3559	0.0170	0.0	0.0
3560	0.0381	0.0	0.0
3561	0.0009	15.9	0.0
3562	0.0014	0.0	0.0

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
3563	0.0027	30.1	0.2
3564	0.0343	0.0	0.0
3568	0.0034	24.9	0.2
3569	0.0028	61.9	0.4
3570	0.0031	120.2	0.8
3571	0.0026	2309.9	13.1
3572	0.0026	894.8	5.2
3573	0.0011	1358.3	3.2
3574	0.0001	2327.5	0.7
3575	0.0000	4968.6	0.4
3576	0.0003	315.1	0.2
3577	0.0002	423.9	0.2
3578	0.0002	2626.9	1.0
3579	0.0014	2021.9	6.3
3580	0.0010	2751.8	6.2
3581	0.0012	2060.7	5.4
3582	0.0006	1587.1	2.1
3583	0.0005	6391.8	7.2
3633	0.0053	0.0	0.0
3634	0.0033	0.0	0.0
3635	0.0033	0.0	0.0
3636	0.0071	0.0	0.0
3637	0.0291	0.7	0.0
3640	0.0023	0.0	0.0
3641	0.0028	0.0	0.0
3642	0.0012	55.8	0.2
3644	0.0001	86.7	0.0
3645	0.0044	2753.3	26.8
3646	0.0014	3186.0	10.0
3647	0.0025	314.1	1.7
3648	0.0013	536.3	1.5
3649	0.0008	734.0	1.3
3650	0.0007	147.8	0.2
3651	0.0001	976.0	0.2
3652	0.0001	4403.5	1.1
3653	0.0005	2432.1	2.6
3654	0.0004	2604.8	2.2
3655	0.0008	2588.4	4.3

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
3656	0.0012	1852.2	4.8
3657	0.0009	465.8	0.9
3658	0.0000	115.2	0.0
3659	0.0000	0.0	0.0
3707	NA	0.0	NA
3708	0.0006	0.0	0.0
3709	0.0012	0.0	0.0
3710	0.0010	0.0	0.0
3711	0.0485	0.0	0.0
3712	0.0281	0.0	0.0
3713	0.0028	0.0	0.0
3717	0.0022	0.0	0.0
3718	0.0026	857.0	4.8
3719	0.0092	2600.7	52.2
3720	0.0026	3112.7	17.9
3721	0.0044	175.2	1.7
3722	0.0008	361.0	0.6
3723	0.0046	159.7	1.6
3724	0.0023	3.8	0.0
3725	0.0018	147.1	0.6
3726	0.0005	490.3	0.5
3727	0.0001	1224.8	0.2
3728	0.0000	1710.1	0.1
3729	0.0004	1375.1	1.1
3730	0.0010	281.3	0.6
3731	0.0011	0.0	0.0
3780	NA	0.0	NA
3783	0.0054	0.0	0.0
3786	0.0046	0.0	0.0
3787	0.0057	0.0	0.0
3789	NA	NA	NA
3791	0.0064	168.7	2.3
3792	0.0021	1303.9	5.9
3793	0.0018	1844.7	7.4
3794	0.0034	107.9	0.8
3795	0.0055	238.6	2.9
3796	0.0034	289.8	2.1
3797	0.0029	376.3	2.4

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
3798	0.0034	1138.0	8.6
3799	0.0031	742.8	5.1
3800	0.0046	99.4	1.0
3801	0.0041	744.2	6.6
3802	0.0014	766.4	2.4
3803	0.0005	62.1	0.1
3804	0.0006	0.0	0.0
3805	0.0020	0.0	0.0
3855	NA	NA	NA
3864	0.0094	161.2	3.3
3865	0.0019	423.8	1.8
3866	0.0019	316.8	1.3
3867	0.0020	492.9	2.2
3868	0.0096	177.7	3.7
3869	0.0049	931.3	9.9
3870	0.0096	0.0	0.0
3871	0.0060	686.5	9.1
3872	0.0084	318.3	5.8
3873	0.0099	326.2	7.1
3874	0.0141	1236.7	38.3
3875	0.0137	2156.8	64.9
3876	0.0068	37.2	0.5
3877	0.0063	0.0	0.0
3878	0.0006	0.0	0.0
3936	0.1930	0.0	0.0
3938	NA	NA	NA
3939	0.0006	183.3	0.2
3940	0.0009	845.6	1.7
3941	0.0022	1397.5	6.6
3944	0.1145	1094.8	274.9
3945	0.0162	793.7	28.2
3946	0.0073	812.2	13.1
3947	0.0129	3622.5	102.1
3948	0.0136	1103.0	33.0
3949	0.0104	66.4	1.5
3950	0.0100	0.0	0.0
4000	NA	0.0	NA
4004	NA	NA	NA

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
4008	NA	0.0	NA
4012	0.0009	151.8	0.3
4014	0.0011	0.0	0.0
4018	0.0100	787.9	17.2
4019	0.0121	590.7	15.6
4020	0.0523	120.8	13.8
4021	0.0080	2371.2	41.4
4022	0.0074	776.1	12.7
4023	0.0039	117.4	1.0
4024	0.0047	0.0	0.0
4081	0.0228	0.0	0.0
4085	0.5632	0.0	0.0
4087	0.1044	237.3	54.3
4088	0.0565	1127.2	139.5
4092	0.0673	392.9	58.0
4093	0.0471	1806.7	186.5
4094	0.0058	1301.3	16.7
4095	0.0093	2507.1	51.0
4096	0.0138	1519.6	45.9
4158	NA	NA	NA
4159	NA	0.0	NA
4160	0.0072	46.7	0.7
4162	0.1151	3091.5	779.9
4163	0.0332	3324.5	241.6
4164	0.0176	2153.5	83.1
4165	0.0400	1811.2	158.9
4166	0.0567	2052.0	255.1
4167	0.0038	1135.8	9.4
4168	0.0050	5486.1	60.7
4169	0.0101	8869.7	197.0
4228	0.0100	0.0	0.0
4229	NA	0.0	NA
4235	0.0885	2049.4	397.4
4236	0.0593	2294.4	298.4
4237	0.0095	3918.1	81.2
4238	0.0184	1181.5	47.8
4240	NA	450.3	NA
4241	0.0130	3907.9	111.4

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
4242	0.0065	2339.7	33.3
4302	0.0082	0.0	0.0
4306	0.0256	415.0	23.3
4307	0.0158	256.7	8.9
4311	0.0211	118.9	5.5
4313	0.0423	70.9	6.6
4314	0.1690	1327.3	491.6
4315	0.0528	2445.4	283.0
4376	NA	3.4	NA
4382	NA	200.0	NA
4383	0.0437	1202.3	115.0
4384	0.0028	95.5	0.6
4385	NA	122.8	NA
4386	NA	241.6	NA
4387	NA	3317.8	NA
4388	NA	3667.0	NA
4389	NA	NA	NA
4442	0.7577	0.0	0.0
4443	NA	0.0	NA
4458	NA	0.0	NA
4459	NA	30.1	NA
4460	NA	58.0	NA
4461	NA	2149.7	NA
4462	NA	332.0	NA
4463	NA	780.2	NA
4516	NA	0.0	NA
4522	NA	0.0	NA
4527	NA	6.6	NA
4528	NA	3.5	NA
4529	NA	0.0	NA
4531	NA	0.0	NA
4532	NA	0.0	NA
4533	NA	0.0	NA
4534	NA	147.7	NA
4535	NA	562.9	NA
4536	0.4018	260.3	229.2
4537	NA	291.4	NA
4543	NA	0.0	NA

Table E-3 (continued).

Grid Cell ID	Methane EF	AF (MMscf/yr)	Methane Emissions (2013) (kg/h)
4590	NA	0.0	NA
4594	NA	0.0	NA
4595	NA	0.0	NA
4597	NA	0.0	NA
4599	NA	0.0	NA
4600	NA	123.8	NA
4605	NA	0.0	NA
4606	NA	0.0	NA
4607	NA	0.0	NA
4608	NA	136.0	NA
4609	0.0101	83.6	1.9
4610	NA	79.8	NA
4672	NA	0.0	NA
4674	NA	NA	NA
4679	NA	0.0	NA
4680	NA	0.0	NA
4682	NA	0.0	NA
4683	0.2531	0.0	0.0
4684	0.0137	0.0	0.0
4753	NA	0.0	NA
4756	NA	0.0	NA
4757	NA	0.0	NA
4758	NA	0.0	NA

E.4 Per grid cell scaling factors

Another way of looking at the scaling of emissions per grid cell, is to calculate a scaling factor; where the ratio of *[Natural Gas Production 2013]/[Natural Gas Production 2009]* is determined for each individual grid cell (instead of a single average ratio for the Barnett Shale Region) and then used to scale 2009 methane emissions for each particular grid cell; which is equivalent to multiplying the emission factor times the activity factor for each grid cell, and then summing up all the emissions. Total estimated methane emissions would be 46,745 kg/h; with the results showing significant spatial differences that cause emissions to grow by a factor of three.

The increase of emissions points to a skewed spatial distribution of production changes from 2009 to 2013 within the Barnett Shale. Production decreases in some sections of the region, while others show increases that are considerably higher than 8%. From the initial (baseline year) dataset, 30% of the grid cells had a 2013 production of zero, a subset of grid cells with non-zero 2013 production is used for the remaining analyses in this section.

Figure E-3 bins the grid cells by the magnitude of the scaling factors (*the ratio of 2013 production divided by 2009 production*), showing the percent of methane emissions (for the baseline year) falling into each scaling factor bin. The map in the right side of Figure E-3 shows the scaling factor of each grid cell, highlighting the heterogeneous shifts in production. Roughly 37% of the emissions occur in grid cells that have a scaling factor between 1.0 and 10, and 6% of the emissions would be multiplied by a scaling factor greater than 10. In terms of grid cells, roughly 10% of them have a scaling factor greater than 10.

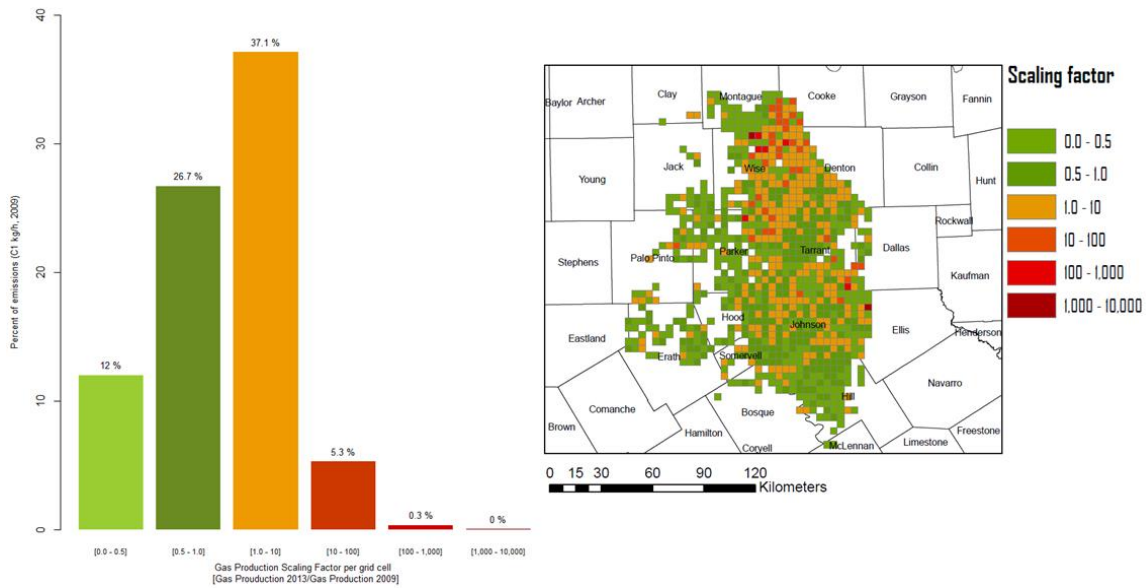


Figure E-3. Distribution of estimated methane emissions for the baseline year, categorized by grid cell scaling factors (*left*), and their spatial distribution (*right*).

E.5 References

- (1) M. Vazquez and H.D. Beggs, Correlations for Fluid Physical Property Prediction, *Journal of Petroleum Technology*, June, 1980, 968-970.
- (2) Allen DT, Torres VM, Thomas J, Sullivan D, Harrison M, et al. 2013. Measurements of Methane Emissions at Natural Gas Production Sites: Study Appendices and Database. Austin: Univ. Tex. <http://dept.ceer.utexas.edu/methane/study/>
- (3) Texas Commission on Environmental Quality, Calculating Volatile Organic Compounds (VOC) Flash Emissions from Crude Oil and Condensate Tanks at Oil and Gas Production Sites, Available at http://www.tceq.texas.gov/assets/public/permitting/air/Guidance/NewSourceReview/guidance_flashemission.pdf (Accessed 6/22/2014).
- (4) Green, D. W., & Perry, R. H. (2007). *Perry's Chemical Engineers' Handbook*. Blacklick, OH: McGraw Hill.

(5) Zavala-Araiza, D., Sullivan, D., & Allen, D. T. (2014). Atmospheric hydrocarbon emissions and concentrations in the Barnett Shale natural gas production region. *Environmental Science & Technology*, 48(9), 5314-5321.

BIBLIOGRAPHY

- Adgate, J. L., Goldstein, B. D., McKenzie, L. M., (2014) Potential Public Health Hazards, Exposures and Health Effects from Unconventional Natural Gas Development. *Env. Sci. Tech*, in press.
- Allen DT, Torres VM, Thomas J, Sullivan D, Harrison M, et al. (2013). Measurements of Methane Emissions at Natural Gas Production Sites: Study Appendices and Database. Austin: Univ. Tex. <http://dept.ceer.utexas.edu/methane/study/>
- Allen, D.T., Torres, V.M., Thomas, J., Sullivan, D., Harrison, M., Hendler, A., Herndon, S.C., Kolb, C.E., Fraser, M., Hill, A.D., Lamb, B.K., Miskimins, J., Sawyer, R.F., and Seinfeld, J.H. Measurements of Methane Emissions at Natural Gas Production Sites in the United States, *Proceedings of the National Academy of Sciences*, 110, 17768-17773, doi: 10.1073/pnas.1304880110 (2013).
- Alvarez RA, Pacala SW, Winebrake JJ, Chameides WL, Hamburg, SP (2012). Greater Focus Needed on Methane Leakage from Natural Gas Infrastructure *Proceedings of the National Academy of Sciences* 109: 6435–6440.
- American Gas Association. (1951). *Fuel Flue Gases*.
- American Petroleum Institute and America's Natural Gas Alliance (API/ANGA) Characterizing Pivotal Sources of Methane Emissions from Natural Gas Production,: Summary and Analyses of API and ANGA Survey Responses, Final Report, updated September, 2012, available at: <http://www.api.org/news-and-media/news/newsitems/2012/oct-2012/~//media/Files/News/2012/12-October/API-ANGA-Survey-Report.pdf>
- API. (2009). *Compendium of Greenhouse Gas Emissions Methodologies for Oil and Natural Gas Industry*. Washington, DC: American Petroleum Institute.

- Argonne National Laboratory. (2011). Life-Cycle Analysis of Shale Gas and Natural Gas . Oak Ridge, TN: U.S. Department of Energy.
- Argonne National Laboratory. (n.d.). The Greenhouse Gases, Regulated Emissions, and Energy Use In Transportation Model, GREET 1.8d.1. Retrieved 6 16, 2014, from GREET: <http://greet.es.anl.gov/>
- Armendariz, A. (2009). Emissions from natural gas production in the Barnett shale area and opportunities for cost-effective improvements. Austin, Texas: Environmental Defense Fund.
- Aydin, M.; Verhulst, K. R.; Saltzman, E. S.; Battle, M. O.; Montzka, S. A.; Blake, D. R.; Tang, Q.; Prather, M. J. Recent decreases in fossil-fuel emissions of ethane and methane derived from firm air. *Nature* 2011, 476 (7359), 198-201.
- Babusiaux D, Pierru A (2007). Modelling and allocation of CO2 emissions in multiproduct industry: The case of oil refining *Applied Energy*, 84: 828-841.
- Brandt AR, Heath GA, Kort EA, O’Sullivan F, Pétron G, Jordaan, SM, Tans P, Wilcox J, Gopstein, AM, Arent D, et al. (2014). Methane Leaks from North American Natural Gas Systems. *Science*, 343: 733-735.
- Caulton D. R., Shespon P. B., Santoro, R. L., Sparks J. P, Howarth, R. W., Ingraffea, A. R., Cambaliza M. O. L., Sweeney, C., Karion, A., Davis, K. J., Stirm, B. H., Montzla S. A., Miller, B. R. (2014). Toward a better understanding and quantification of methane emissions from shale gas development. *Proceedings of the National Academy of Sciences*, 111, 6237 – 6242.
- Cimorelli, A. J.; Perry, S. G.; Venkatram, A.; Weil, J. C.; Paine, R. J.; Wilson, R. B.; Lee, R. F.; Peters, W. D.; Brode, R W.; Paumier, J. O. (2004) AERMOD: Description of the Model Formulation, EPA, North Carolina.

- Davies, R., Foulger, G., Bindley, A., & Styles, P. (2013). Induced seismicity and hydraulic fracturing for the recovery of hydrocarbons. *Marine and Petroleum Geology*, 45, 171-185.
- East. Res. Group/Sage Environ. Consult. (2011). City of Fort Worth natural gas air quality study: final report, July 13, Fort Worth, TX. Accessed Jan. 2013. http://fortworthtexas.gov/uploadedFiles/Gas_Wells/AirQualityStudy_final.pdf
- Edwards, P.M., Young, C. J., Aikin, K., deGow, J., Dube, W. P., Geiger, F., Gilman, J., Helmig D., Holloway, J. S., Kercher, J., Lerner, B., Martin, R., McLaren R., Parrish, D. D., Peischl J., Roberts, J.M., Ryerson T. B., Thornton, J., Warneke C., Williams, E. J., Brown, S., S., (2013) Ozone photochemistry in an oil and natural gas extraction region during winter: simulations of a snow-free season in the Uintah Basin, Utah, *Atmos. Chem. Phys.*, 13, 9955-8971.
- Efron B.; Tibshirani, R. *An Introduction to the Bootstrap*, Chapman & Hall, CRC Press, (1993).
- EIA, (2013). What is shale gas and why is it important? Retrieved 9/25/2013 from http://www.eia.gov/energy_in_brief/article/about_shale_gas.cfm
- EIA, Monthly Natural Gas Gross Production Report; Retrieved 9/13/2012 from http://www.eia.gov/oil_gas/natural_gas/data_publications/eia914/eia914.html
- EIA. (2011). Glossary. Retrieved 8 12, 2013, from <http://www.eia.gov/naturalgas/annual/pdf/glossary.pdf>
- EIA. (2012). Annual Energy Outlook. DOE/EIA.
- EIA. (2013). Annual Energy Outlook 2013. Retrieved 10 19, 2013, from http://www.eia.gov/energy_in_brief/data/shaleplay_production_AEO2013ER.xlsx
- EIA. (2013). Oil and Gas Supply Module of the National Energy Modeling System: Model Documentation 2013, Appendix 2.c. Retrieved 1, 14, 2014, from [http://www.eia.gov/forecasts/aeo/nems/documentation/ogsm/pdf/m063\(2013\).pdf](http://www.eia.gov/forecasts/aeo/nems/documentation/ogsm/pdf/m063(2013).pdf)

- EIA. (2013). Technically Recoverable Shale Oil and Shale Gas Resources: An Assessment of 137 Shale Formations in 41 Countries Outside the United States. Retrieved June 13, 2013, from <http://www.eia.gov/analysis/studies/worldshalegas/>
- EIA. (2013). The Natural Gas Production, Transmission and Distribution System. Retrieved 5 29, 2013, from <http://www.window.state.tx.us/specialrpt/energy/nonrenewable/gas.php>
- EIA. (2014). Annual Energy Outlook. Washington DC: Department of Energy.
- Ekvall T, Tillman A-M (1997). Open-Loop Recycling: Criteria for Allocation Procedures. The International Journal of Life Cycle Assessment, 2 (3): 155-162.
- Ellsworth, W. L. (2013). Injection-Induced Earthquakes. Science, 341.
- Environmental Protection Agency (EPA), Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2011. April 12, 2013 Retrieved 4/24/2013 from <http://www.epa.gov/climatechange/ghgemissions/usinventoryreport.html>
- Environmental Protection Agency (EPA). Inventory of U.S. Greenhouse Gas Emissions and Sinks. <http://www.epa.gov/climatechange/Downloads/ghgemissions/US-GHG-Inventory-2014-Main-Text.pdf> (accessed 21 June 2014).
- EPA. (2011). Plan to Study the Potential Impacts of Hydraulic Fracturing on Drinking Water Resources. Retrieved 8 15, 2012, from http://www.epa.gov/hfstudy/HF_Study__Plan_110211_FINAL_508.pdf
- EPA. (2013, April 12). Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2011. Retrieved 4 22, 2013, from <http://www.epa.gov/climatechange/ghgemissions/usinventoryreport.html>
- EPA. (2014, April 15). Inventory of U.S. Greenhouse Gas Emissions and Sinks: 1990-2012. Retrieved 6 20, 2014, from <http://www.epa.gov/climatechange/ghgemissions/usinventoryreport.html>

- Gilman, J. B., Lerner B. M., Kuster, W.C., de Gouw, J. A. (2013) Source Signature of Volatile Organic Compounds from Oil and Natural Gas Operations in the Northeastern Colorado. *Environmental Science and Technology*, 47, 1297-1305.
- Green, D. W., & Perry, R. H. (2007). *Perry's Chemical Engineers' Handbook*. Blacklick, OH: McGraw Hill.
- GREET, developed by Argonne National Laboratory, from <http://greet.es.anl.gov/>
- Harrison, M.R.; Shires, T.M.; Wessels, J.K.; Cowgill, R. M; Methane Emissions from the Natural Gas Industry, Volumes 1 – 15, Final Report, GRI-94/0257 and EPA-600/R-96-080, Gas Research Institute and US Environmental Protection Agency, June 1996.
- Harrison, M.R.; Shires, T.M.; Wessels, J.K.; Cowgill, R.M.; Methane Emissions from the Natural Gas Industry, EPA/600/SR-96/080 June 1997.
- Herndon SC, Floerchinger C, Roscioli JR, Yacovitch TI, Franklin JF, et al. (2013). Measuring methane emissions from industrial and waste processing sites using the dual tracer flux ratio method. Presented at Annu. Meet. Am. Geophys. Union, Dec. 2013, San Francisco
- Herndon SC, Floerchinger C, Roscioli JR, Yacovitch TI, Franklin JF, Shorter JH, Kolb, CE, Subramanian R, Robinson AL, Molina LT, et al. (2013) Measuring methane emissions from industrial and waste processing sites using the dual tracer flux ratio method Annual Meeting the American Geophysical Union, San Francisco, December 2013.
- Herndon SC, Jayne JT, Zahniser MS, Worsnop DR, Knighton B, Alwine E, Lamb BK, Zavala M, Nelson DD, McManus JB, et al. (2005) Characterization of urban pollutant emission fluxes and ambient concentration distributions using a mobile laboratory with rapid response instrumentation *Faraday Discuss.* 130: 327–339.
- Howarth, R. W., Santoro, R., & Ingraffea, A. (2011). Methane and the greenhouse-gas footprint of natural gas from shale formations. *Climatic Change*, 106, 679-690.

- ICF. (2014). Economic Analysis of Methane Emission Reduction Opportunities in the U.S. Onshore Oil and Natural Gas Industries. Prepared for Environmental Defense Fund. Available at http://www.edf.org/sites/default/files/methane_cost_curve_report.pdf (Retrieved 7/3/2014).
- IEA. (2012). World Energy Outlook. Retrieved 5 29, 2013, from <http://www.worldenergyoutlook.org/publications/weo-2012/>
- IPCC. (2007). Fourth Assessment Report (AR4), Climate Change 2007: The Physical Science Basis.
- IPCC. (2013). Fifth Assessment Report (AR5), Climate Change 2013: The Physical Science Basis. Retrieved 10 19, 2013, from http://www.climatechange2013.org/images/uploads/WGIAR5_WGI-12Doc2b_FinalDraft_All.pdf
- Jaramillo P, Griffin WM, Matthews HS (2007). Comparative lifecycle air emissions of coal, domestic natural gas, LNG, and SNG for electricity generation. *Environ. Sci. Technol.* 41: 6290–6296.
- Jenner, S., & Lamadrid, A. J. (2013). Shale gas vs. coal: Policy implications from environmental impact comparisons of shale gas, conventional gas, and coal on air, water, and land in the United States. *Energy Policy*, 53, 442-453.
- Jimenez, R., Herndon, D., Shorter, J., Nelson, D., McManus, J., Zahniser, M., (2005). Atmospheric trace gas measurements using dual quantum cascade laser mid-infrared absorption spectrometer. *Novel In-Pane Semiconductor Lasers IV*. Bellingham SPIE-INT Society Optical Engineering. Proc. SPIE 5738, 318 – 331.
- Karion A, Sweeney C, P'etron G, Frost G, Hardesty RM, et al. (2013). Methane emissions estimate from airborne measurements over a western United States natural gas field. *Geophys. Res. Lett.* 40:4393–97

- Katzenstein AS, Doezema LA, Simpson IJ, Blake DR, Rowland FS (2003) Extensive regional atmospheric hydrocarbon pollution in the southwestern United States. *Proc. Natl. Acad. Sci. U.S.A.* 100: 11975–11979.
- Kirchgesner, D.A.; Lott, R.A. Cowgill, R.M.; Harrison, M.R.; Shires, T.M. Estimate of Methane Emissions from the U.S. Natural Gas Industry, *Chemosphere*, 35, 1365-1390, 1997.
- Kolb CE, Herndon SC, McManus JB, Shorter JH, Zahniser MS, et al. (2004). Mobile laboratory with rapid response instruments for real-time measurements of urban and regional trace gas and particulate distributions and emission source characteristics. *Environ. Sci. Technol.* 38:5694–703
- Konschnik, K., (2014). Shale Gas Development: A Smart Regulation Framework. *Environmental Science & Technology*.
- Kort, E. A.; Eluszkiewicz, J.; Stephens, B.; Miller, J.; Gerbig, C.; Nehrkorn, T.; Daube, B.; Kaplan, J.; Houweling, S.; Wofsy, S. Emissions of CH₄ and N₂O over the United States and Canada based on a receptor-oriented modeling framework and COBRA-NA atmospheric observations. *Geophys. Res. Lett.* 2008, 35, L18808.
- Lamb BK, Shorter JH, McManus JB, Kolb CE, Mosher BW, et al. (1995). Development of atmosphere tracer methods to measure methane emissions from natural gas facilities and urban areas. *Environ. Sci. Technol.* 29:1468–78
- Laurenzi IJ, Jersey GR (2013). Life Cycle Greenhouse Gas Emissions and Freshwater Consumption of Marcellus Shale Gas *Environ. Sci. Tech.* 47: 4896-4903.
- Lozano Maya, J. (2013). The United States experience as a reference of success for shale gas development: The case of Mexico. *Energy Policy*, 70-78.
- M. Vazquez and H.D. Beggs, Correlations for Fluid Physical Property Prediction, *Journal of Petroleum Technology*, June, 1980, 968-970.

- MacKay, D. J., & Stone, T. J. (2013). Potential Greenhouse Gas Emissions Associated with Shale Gas Extraction and Use. London: DECC.
- McGaughey, G., Sullivan, D., Smith, D., McDonald-Buller, E., Allen, D. (2009) Meteorological Conditions and Associated Temporal Trends in Total Non-Methane Hydrocarbon and Benzene Concentrations in the Corpus Christi Area. The University of Texas at Austin. Available at http://www.utexas.edu/research/ceer/ccaqp/PDF/CCNATReports/Part3_CCNAT_FY2008_2009_Air_Toxics_Conceptual_Model.pdf (Retrieved 7/3/2014)
- Medlock, K. B. (2012) Modeling the implications of expanded US shale gas production. *Energy Strategy Reviews*, 1, 33-41.
- Melikoglu, M. (2014). Shale Gas: Analysis of its role in the global market. *Renewable and Sustainable Energy Reviews*, 460-468.
- Miller SM, Wofsy SC, Michalak AM, Kort EA, Andrews AE, Biraud SC, Dlugokencky EJ, Eluszkiewicz J, Fischer ML, Janssens-Maenhout G, et al. (2013) Anthropogenic emissions of methane in the United States. *Proc. Natl. Acad. Sci. U.S.A*, 110, 20018–20022.
- Nelson, D., Shorter, J., McManus, J., Zahniser, M., (2006) Sub-par-per-billion detection of nitric oxide in air using thermoelectrically cooled mid-infrared quantum cascade laser spectrometer. *Applied Physics B-Lasers and Optics* 75, 343 – 350.
- NETL. (2011). Life Cycle Greenhouse Gas Inventory of Natural Gas Extraction, Delivery and Electricity Production. U.S. Department of Energy.
- NETL. (2012). Role of Alternative Energy Sources: Natural Gas Technology Assessment. U.S. Department of Energy.
- Nicot, J., & Scanlon, B. R. (2012). Water Use for Shale-Gas Production in Texas. *Environmental Science & Technology*, 46, 3580-3586.

- Osborn, S. G., Vengosh, A., Warner, N. R., & Jackson, R. B. (2011). Methane contamination of drinking water accompanying gas-well drilling and hydraulic fracturing. *Proceedings of the National Academy of Science*, 108, 8172-8176.
- O'Sullivan, F., & Paltsev, S. (2012). Shale gas production: potential versus actual greenhouse gas emissions. *Environmental Research Letters*, 7, 1-7.
- Pacsi AP, Alhajeri NS, Zavala-Araiza D, Webster MD, Allen DT (2013). Regional air quality impacts of increased natural gas production and use in Texas. *Environ. Sci. Tech.* 47: 3521-3527.
- Peischl J, Ryerson TB, Brioude J, Aikin KC, Andrews AE, Atlas E, Blake D, Daube BC, de Gouw JA, Dlugokencky E, et al. (2013) Quantifying Sources of Methane Using Light Alkanes in the Los Angeles Basin, California *Journal of Geophysical Research Atmospheres* 2013, 118:4974-4990.
- Perry, S.G.; Cimorelli, A. J.; Paine, R. J.; Brode, R W.; Weil, J. C.; Venkatram, A.; Wilson, R. B.; Lee, R. F.; R. F. (2005) AERMOD: A Dispersion Model for Industrial Source Applications. Part II: Model Performance against 17 Field Study Databases; *Journal of Applied Meteorology*, 44, 682-693.
- Petron G, Frost G, Miller BR, Hirsch AI, Montzka SA, Karion A, Trainer M, Sweeney C, Andrews AE, Miller L, et al. (2012). Hydrocarbon Emissions Characterization in the Colorado Front Range: A Pilot Study *J. Geophys. Res.*, 117(D4): D04304.
- Pétron, G.; Karion, A.; Sweeney, C.; Miller, B.; Montzka, S.; Frost, G.; Trainer, M.; Tans, P.; Andrews, A.; Kofler, J.; Helmig, D.; Guenther, D.; Dlugokencky, E.; Lang, P.; Newberger, T.; Wolter, S.; Hall, B.; Novelli, P.; Brewer, A.; Conley, S.; Hardesty, M.; Banta, R.; White, A.; Noone, D.; Wolfe, D.; Schnell, R. (2014) A new look at methane and nonmethane hydrocarbon emissions from oil and natural gas operations in the Colorado Denver –Julesburg Basin. *J. Geophys. Res. Atmos.* 119, 1-17.

- Prasino Group, Final Report for determining bleed rates for pneumatic devices in British Columbia. Report to British Columbia Ministry of Environment, December 2013.
- Rahm, B., & Riha, S. (2012). Toward strategic management of shale gas development: Regional, collective impacts on water resources. *Environmental Science & Policy*, 17, 12-23.
- Rahm, D. (2011). Regulating hydraulic fracturing in the shale gas plays: The case of Texas. *Energy Policy*, 39, 2974-2981.
- Rail Road Commission of Texas . (2012). Newark, East (Barnett Shale) Statistics. Retrieved 12 18, 2012, from <http://www.rrc.state.tx.us/data/fielddata/barnettshale.pdf>
- Rail Road Commission of Texas, Texas Gas Well Gas Production in the Newark, East (Barnett Shale) Field – 1993-2011. Retrieved 9/11/2012 from http://www.rrc.state.tx.us/barnettshale/NewarkEastField_1993-2011.pdf
- Richardson, N., Gottlieb, M., Krupnick., Wiseman, H., (2013). The State of State Shale Gas Regulation. RFF Report, Washington DC.
- Roy A. A., Adams, P. J., Robinson A. L., (2014) Air pollutant emissions from the development, production, and processing of Marcellus Shale natural gas. *Journal of the Air & Waste Management Association*, 64, 19-37.
- SENER (Department of Energy, Mexico). (n.d.). Why is Shale Gas/Oil and why is it important? Retrieved 6 2014, from http://www.energia.gob.mx/webSener/shale/shale_en.html
- Shorter JH, McManus JB, Kolb CE, Allwine EJ, Siverson R, et al. (1997). Collection of leakage statistics in the natural gas system by tracer methods. *Environ. Sci. Technol.* 31:2012–19
- Texas Commission on Environmental Quality (TCEQ). Barnett Shale Special Emissions Inventory Phase I. 2011. <https://www.tceq.texas.gov/assets/public/implementation/air/ie/pseiforms/Barnett%20Shale%20Area%20Special%20Inventory.pdf> (accessed 21 June 2014).

- Texas Commission on environmental Quality, Air Monitoring Sites, Retrieved 9/11/2012 from http://www.tceq.texas.gov/airquality/monops/sites/mon_sites.html
- Texas Commission on environmental Quality, Barnett Shale Geological Area; Retrieved 9/11/2012 from <http://www.tceq.texas.gov/airquality/barnettshale>
- Texas Commission on environmental Quality, Barnett Shale Special Inventory, Phase Two Workbook (Excel), Retrieved 10/7/2012 from <http://www.tceq.texas.gov/assets/public/implementation/air/ie/pseiforms/bshaleworkbook.xls>
- Texas Commission on Environmental Quality, Calculating Volatile Organic Compounds (VOC) Flash Emissions from Crude Oil and Condensate Tanks at Oil and Gas Production Sites, Available at http://www.tceq.texas.gov/assets/public/permitting/air/Guidance/NewSourceReview/guidance_flashemission.pdf (Accessed 6/22/2014).
- Texas Commission on environmental Quality, Dallas-Fort Worth Eight-Hour Ozone SIP Modeling (2006 Episode), Available at <http://www.tceq.state.tx.us/airquality/airmod/data/dfw8h2> (accessed 6/22/2014)
- Texas Commission on environmental Quality, Point Source Emissions inventory, Retrieved 9/19/2012 from <http://www.tceq.texas.gov/airquality/point-source-ei/psei.html#barnett2>
- U.S. Environmental Protection Agency (EPA), Office of Air Quality Planning and Standards. Oil and Natural Gas Sector Pneumatic Devices; Report for Review Panel, April 2014.
- U.S. Environmental Protection Agency (EPA). Greenhouse Gas Reporting Program, Data for Reporting Year 2012, 2013.
- U.S. Environmental Protection Agency, Natural Gas STAR Program. Available at <http://epa.gov/gasstar/> (Retrieved 7/3/2014)

- Venkatesh A, Jaramillo P, Griffin, WM, Matthews HS (2012). Implications of changing natural gas prices in the United States electricity sector for SO₂, NO_x, and life cycle of GHG emissions *Environ. Res. Lett.* 7: 034018.
- Wang, M., Hanjie, L., & Molburg, J. (2004). Allocation of Energy Use in Petroleum Refineries to Petroleum Products. *The International Journal of Life Cycle Assessment*, 9 (1), 34-44.
- Wecht KJ, Jacob DJ, Sulpizio MP, Santoni GW, Wofsy SC, Parker R, Bosch H. and Worden J. (2014). Spatially resolving methane emissions in California: constraints from the CalNex aircraft campaign and from present (GOSAT, TES) and future (TROPOMI, geostationary) satellite observations. *APCD*, 14: 4119-4148.
- Wigley, T. L. (2011). Coal to gas: the influence of methane leakage. *Climatic Change*, 108, 601-608.
- Yacovitch, T., Herndon S., Roscioli, J., Floerchinger C., et al., (2014). Demonstration of an Ethane Spectrometer for Methane Source Identification. *Environmental Science and Technology* 48, 8028-8034.
- Yaping, X.; Logan, J.; Jacob, D.; Hudman, R.; Yantosca, R.; Blake, D. (2008) Global budget of ethane and regional constraints on U.S. sources. *J. Geophys Res.* 113, D21306.
- Zavala-Araiza, D., Sullivan, D., & Allen, D. T. (2014). Atmospheric hydrocarbon emissions and concentrations in the Barnett Shale natural gas production region. *Environmental Science & Technology*, 48(9), 5314-5321.