

Multi-scale Measurements of Greenhouse Gas Emissions at U.S. Natural Gas Liquefaction Terminals

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Abstract:

Addressing methane emissions across the liquefied natural gas (LNG) supply chain is key to reducing climate impacts of LNG. Actions to address methane emissions have emphasized the importance of the use of measurement-informed emissions inventories, given the systematic underestimation in official GHG emission inventories. Despite significant progress in field measurements of GHG emissions across the natural gas supply chain, no detailed measurements at US liquefaction terminals are publicly available. In this work, we conduct multiscale, periodic measurements of methane and carbon dioxide emissions at two US LNG terminals over a 16-month campaign. We find that methane emissions intensity varied from 0.007% to 0.045%, normalized to methane in LNG production. Carbon dioxide emissions accounted for over 95% of total GHG emissions using 100-year global warming potential (GWP) for methane. Thus, contrary to observations across other natural gas supply chain segments, we find that reported GHG emissions intensity closely matches measurement informed GHG emissions intensity of 0.24 – 0.27 kg CO₂e/kg CH₄. In the context of developing LNG supply chain emissions intensity, we conclude that the use of Greenhouse Gas Reporting Program emissions intensity provides reasonably accurate estimates of total GHG emissions at LNG terminals.

Synopsis:

This study presents results from the first multi-scale measurements of GHG emissions at two US liquefaction terminals. We find measured GHG emissions intensity to be consistent with reported intensities.

Keywords: Methane emissions, Measurement-informed inventory, LNG, Supply chain, Climate impacts

1 Introduction

Global demand for liquefied natural gas (LNG) is expected to grow in the future, with US LNG exports to grow by 18% by 2025.^{1,2} This has raised significant concerns about its climate impact, and further underscored by increased demand for non-Russian sources of natural gas in Europe.³⁻⁶ The US Department of Energy (DoE) has temporarily paused review of new LNG export applications in part to update prior analysis of the climate impacts of US LNG.⁷ With a global warming potential 84 times that of carbon dioxide over a 20-year period, methane emissions along the LNG supply chain are a key component of climate impacts of LNG.⁸ Recent research has demonstrated that methane leakage over 4% threatens to erode the climate benefits of natural gas over coal.^{9,10} Thus, addressing methane emissions across the natural gas supply chain is a key component of near-term action to reduce climate impacts of LNG.¹¹

Measurements of methane emissions across the oil and gas supply chain have identified systematic underestimation in official GHG emissions inventories.¹²⁻¹⁵ Thus, global efforts to address methane emissions from the LNG supply chain have emphasized the importance of the use of measurement-informed emissions inventories.¹⁶⁻¹⁹ Furthermore, without a measurement-based and verifiable emissions estimates, it would be impossible to track progress towards national, global, or corporate mitigation targets.¹⁸ The European Commission finalized methane regulations that set measurement and reporting requirements for natural gas and LNG imported into the EU market.²⁰ Several voluntary initiatives, including the US DoE's Measurement, Monitoring, Reporting, and Verification (MMRV) framework and the Oil & Gas Methane Partnership (OGMP) 2.0, propose to establish accurate and transparent emission reporting frameworks for oil and gas suppliers.^{21,22}

In the US, recent measurement campaigns have seen broad deployment of new technologies and highlighted the importance of accurately estimating the frequency and duration of intermittent emission events.^{23–25} Multi-scale measurements of methane emissions at upstream facilities have enabled emissions characterization across a wide range of spatial and temporal scales.^{23,25–28} While a disproportionate number of field campaigns have estimated emissions from the upstream production segment, recent measurements have also focused on midstream facilities. These midstream campaigns have been used to update state and national emissions inventories, identify challenges in measuring emissions from complex facilities.^{29–34}

Despite significant expansion in direct measurements of methane emissions across the LNG supply chain, no direct measurements at US liquefaction terminals are publicly available. This represents a key gap in developing measurement-informed LNG supply chain emissions inventories. Measurements at liquefaction terminals are unique compared to other segments of the supply chains. Liquefaction terminals are significantly larger in size with multi-level structures that have emissions sources spread across multiple levels resulting in complex plume dynamics. This makes ground-based measurements challenging because multiple emissions sources are likely to be inaccessible.³⁵ Furthermore, the physical size and critical security interests of the facility pose additional safety and logistical challenges in conducting measurements compared to upstream or midstream facilities. Outside the US, there was one recent ground-based measurement campaign at liquefaction terminals.³⁶ This study deployed a differential absorption LIDAR instrument from a mobile platform, limiting its ability to quantify emissions from inaccessible locations.

In this study, we present results from the first, multi-scale periodic surveys of GHG emissions at Sabine Pass (Louisiana) and Corpus Christi (Texas) liquefaction terminals over 16 months. By

tracking methane emissions from each source, we demonstrate the role of high-resolution aerial surveys in informing follow-up maintenance activities to reduce methane emissions. By presenting measurement informed GHG emissions intensities at US LNG terminals, this work enables the development of a measurement informed GHG emissions intensity of US LNG supply chains.

2 Methods

2.1 Field Measurements

Multi-scale methane emission measurements were conducted at the Sabine Pass and Corpus Christi liquefaction terminals in Louisiana and Texas, respectively, which together accounted for 51% of US LNG exports in 2023.

The liquefaction terminal measurements followed recent protocols established at production and midstream facilities and was divided into three phases: baseline phase to develop an initial snapshot estimate of whole-site CH₄ and CO₂ emissions, an enhanced monitoring (EM) phase that involved a series of periodic measurements over 6-10 months at each liquefaction facility, and an end-of-project (EOP) verification phase.^{23,24,29,30}

Three technologies were used to detect and quantify methane and carbon dioxide emissions at the liquefaction terminals, including an aerial LIDAR plume identification system by Bridger Photonics (Bridger), an aerial mass-balance measurement using cavity ring-down spectroscopy by ChampionX (ChampionX), and a ground-based Optical Gas Imaging (OGI) camera survey (SI section S1).

Bridger scanned each liquefaction facility with multiple passes per scan and completed several scans during a survey within 1-2 days. A pass refers to a one time North-South or South-North swath, and each scan is a collection of individual Bridger passes over the site. The average emission rate across all Bridger passes in one survey of each emission source was calculated and then aggregated across all sources to obtain as-measured site-level emission rate. Missing quantification estimates for individual passes were assigned rates following standard procedures (SI sections S4). In total, Bridger conducted 49 scans in 13 surveys at the two facilities as part of the campaign.

ChampionX used a mass balance approach to estimate whole-site methane and carbon dioxide emissions. We conducted one successful measurement flight at both site 1 and site 2 in the baseline phase. In the EOP, there were four successful measurement flights for site 1 and two for site 2 (SI section S2).

Ground-based OGI surveys detected emissions using a FLIR GF-320 infrared camera and quantified using a Bacharach Hi-Flow Sampler. Due to thermal interference on OGI camera detection capability and safety requirements on surveyors, OGI cameras may not access equipment such as flare stacks, tanks, and loading arms in the liquefaction terminals. In our study, we used OGI survey as a follow-up at Bridger-identified emission sources.

Across the three phases of measurements, 13 independent aerial surveys were conducted across both sites (SI section S2). Records of maintenance activities between consecutive surveys were provided to the study team for analysis.

In addition, satellite imagery assessments were conducted by Kayrros at both liquefaction terminals using the Sentinel-2 satellite – no emissions were detected over the 16-month period.

2.2 Emissions Inventory Estimates

Two types of bottom-up emissions inventories were estimated for each facility. First, the operator calculated the emissions inventory for each site for the duration of each measurement (called the 'operator-estimated inventory' or OEI) – these included both CH₄ and CO₂ emissions, data from stack tests, ground-based LDAR records, flow rates, and other information necessary to calculate a whole-site emissions inventory (SI section S1). Second, the operator also provided the prior year emissions inventory as submitted to the US EPA through the greenhouse gas reporting program (GHGRP).³⁷

The measurement informed inventory (MII) is calculated based on whole-site emissions estimates incorporating aerial survey data. Time-averaged MII for CH₄ represents the average of MII at each survey through baseline, enhanced monitoring, and EOP phases. Time averaged MII for CO₂ represents the average of MII at baseline and EOP phases, as the EM phase did not include CO₂ measurements. In this work, the MII for methane is based on Bridger's measurements while the MII for CO₂ was based on ChampionX measurements. Results are reported as methane emissions intensity (EI), CO₂ EI, and GHG EI using both 20- and 100-year global warming potential values (SI section S3).

2.3 Quantification Uncertainty

In our study, we propose a new method to estimate the uncertainty in MII that accounts for the skewed estimation error distribution of Bridger's quantification using a quantile regression analysis (SI section S5). The quantile regression analysis provides a conditional distribution of estimation error as a function of the Bridger-estimated emission rate. Nested Monte Carlo (MC) simulations over each pass and source provide uncertainty around whole site MII.

2.4. Treatment of Intermittency

The intermittency observed in aerial surveys may be an artifact of the complex three-dimensional structure of the facility that could prevent full plume development. Thus, a persistent source may appear intermittent to an aerial survey. In this study, we made the conservative assumption that all detected emissions are persistent, irrespective of whether Bridger detected them in all passes.

3 Results

In our study, each survey is considered an independent snapshot estimate of emissions and the emission variation across different surveys is investigated for each facility. We first describe the site-level GHG EI based on the 16-month, multi-scale measurement campaign. We then analyze methane emission variation observed at LNG terminals informed by Bridger measurements, including source-level attribution. We conclude with a discussion of the implication of these data on the development of a global, measurement-informed LNG supply chain EI.

3.1 Site-level GHG emission intensity

Figure 1 shows the CO₂, CH₄, and total GHG (CO₂e) EI across the two liquefaction terminals for the baseline, EM, and the EOP phases. We make three key observations.

First, the CO₂ EI (Figure 1(a,b)) is consistent across surveys at both site 1 and site 2 with average intensities of 0.232 kg CO₂/kg CH₄ and 0.210 kg CO₂/kg CH₄, respectively. Furthermore, the CO₂ EI between the two sites is comparable. The CO₂ emissions from OEI (blue bars) are also consistent with ChampionX mass-balance CO₂ measurements (blue diamonds) because CO₂ emissions are largely from combustion processes and can be accurately estimated using fuel consumption volumes and combustion emissions factors. The higher measured CO₂ EI versus

OEI estimates in site 2 EOP is likely influenced by unfavorable weather conditions during measurements. Specifically, a low cloud base prevented the airplane from flying over the top of the plume, requiring the use of extrapolation to infer the vertical profile.

Second, methane EI varies over 6-fold across surveys ranging from 0.007% to 0.045%. The methane EIs at EM-1 and EM-5 in site 1 are higher than in other surveys and we observe a decreasing trend in methane emission intensity at site 2. The ChampionX mass-balance CH₄ measurements (Figure 1(c,d)) are higher than Bridger as-measured methane emissions and is likely an overestimation. The high density of oil and gas infrastructure around the liquefaction terminals prevented ChampionX from obtaining a clean methane emissions signal associated only with the liquefaction terminals. For example, limitations in the flight path led to the inclusion of sources such as compressor stations, tankers, and wetlands, and other non-LNG terminal sources of methane emissions, resulting in a higher estimate compared to Bridger measurements.

Third, the whole-site GHG emissions intensity (Figure 1(e,f)) remains consistent across measurements as total GHG emission is dominated by CO₂ emissions at liquefaction terminals. CO₂ emission accounts for over 95% of total GHG emission (GWP-100 basis) in each measurement for both sites – thus, variation in methane emissions do not significantly affect the whole-site GHG EI.

Because of the skewed error distribution associated with Bridger's quantification as identified in recent controlled release tests³⁸, the uncertainty range in the methane EI (Figure 1(c,d)) does not follow a normal distribution. This results in an asymmetric confidence interval around the median methane EI, suggesting a small non-zero bias in quantification estimates (SI Section S5).

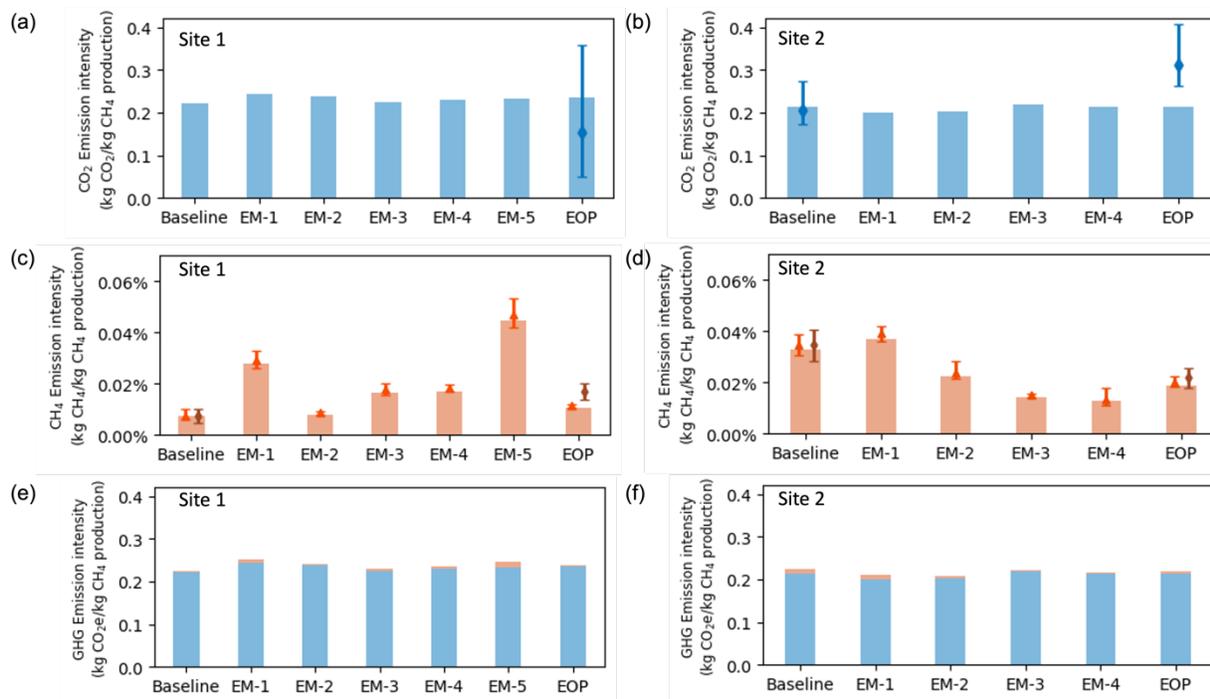


Figure 1. Carbon dioxide and methane emission intensities normalized to natural gas production across two liquefaction terminals. (a,b) CO₂ EI at site 1 and site 2 across baseline, EM, and EOP phases. The CO₂ EI are obtained from OEI (blue bars) and ChampionX measurements (blue diamonds). The ChampionX estimated CO₂ emissions at site 1 baseline is removed because it did not pass quality checks during analysis. (c,d) Bridger as-measured CH₄ EI (orange bars) across all baseline, EM, and EOP phases. The median and 95% bootstrapped confidence interval of Bridger measured methane emission intensity after incorporating Bridger's quantification error (orange triangle) are shown across all measurements. The CH₄ EI with uncertainty range obtained from ChampionX measurements are represented by a brown diamond. (e,f) The GHG emission intensity (GWP-100) at site 1 and site 2 calculated based on CO₂ emissions from OEI and Bridger as-measured CH₄ emission (see SI Section S8 for GWP-20 estimates and SI Section S9 for LNG production normalized EI).

3.2 Role of Maintenance Activities in Reducing Methane Emissions

Figure 2 shows the site-level methane EI across all surveys for the two terminals disaggregated by major source categories. The data for each survey are spaced along the x-axis based on the time of measurement. The vertical lines represent various maintenance activities undertaken over the same time period. We observe significant methane emission reduction from each source after corresponding maintenance activities at both sites. The liquefaction train emissions exhibit a decreasing trend after maintenance, with the lowest liquefaction train emissions in EM-3 being 69% lower than that in EM-1. We conclude that high-resolution aerial surveys can enable timely maintenance for emissions mitigation, especially when aerially detected sources are typically not surveyed by OGI due to inaccessibility.

We also observe significant methane EI variations at both site-level and source-level across surveys at both facilities. At site 1, the methane EI ranges from over 0.04% to below 0.01%. At site 2, the site-level EI shows a decreasing trend over the 16-month campaign, attributable to decreasing liquefaction train emissions after maintenance activities.

The large site-level emission variation in site 1 is caused by the significant variation in flare-related emissions. However, we recommend caution in interpreting flare-related, aerial measurements at liquefaction terminals as the effectiveness of aerial quantification of flare emissions at liquefaction terminals is an open research question (see SI section S7).

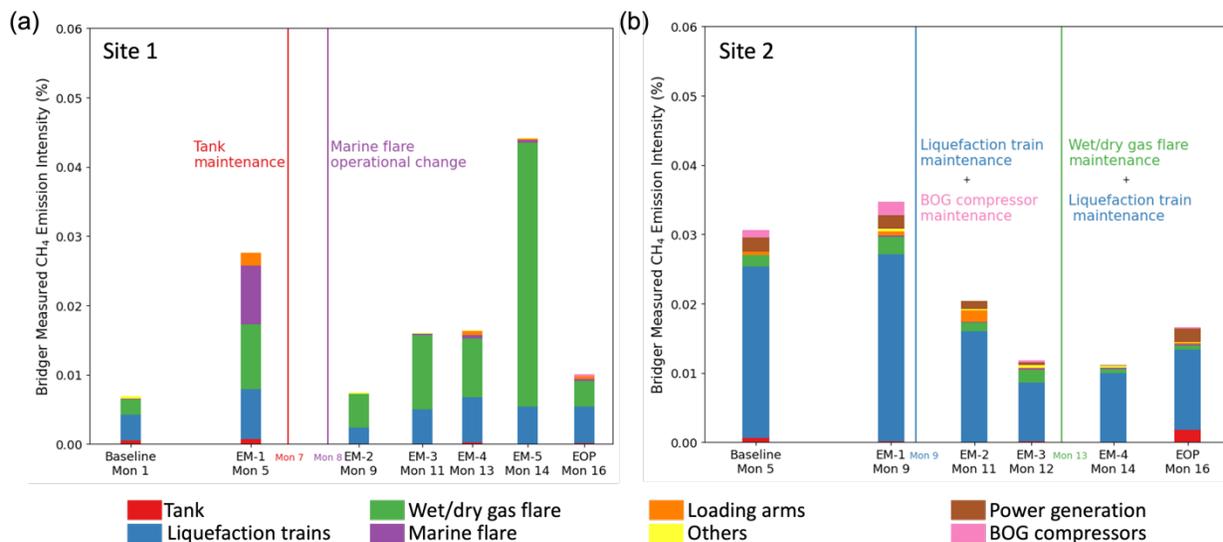


Figure 2. Site-level methane EI across all measurements for two liquefaction terminals ('mon' refers to month), disaggregated by seven major facility groups: tank, liquefaction trains, wet/dry gas flare, marine flare, loading arms, power generation, and BOG compressors. The vertical lines represent different kinds of maintenance operations conducted across these two sites. The specific type of emission reduction is observed after each maintenance.

3.3 Value of OGI Follow-up Survey

Figure 3 shows a reconciliation between the ground based OGI follow-up survey and the aerial Bridger survey. There are four possible scenarios: (1) *OGI identified*: the location of the Bridger-identified emission source is confirmed by the OGI crew; (2) *OGI not found*: the OGI crew did not find any emissions associated with a Bridger-identified emission source; (3) *OGI inaccessible*: the OGI team cannot access a Bridger detected emission source because of safety or logistical considerations; (4) *No OGI follow up*: no OGI follow-up was initiated due to either known limitations of OGI or OGI follow-up would not provide new information for that source. All EI contribution shown in Figure 3 is based on Bridger measurements since the OGI crews did not quantify emissions for all detected sources.

First, OGI cannot be used to estimate whole-site emissions at liquefaction terminals because OGI cameras cannot reach all potential emissions sources at these facilities. At site 1, more than 58% of the Bridger-identified emissions across the whole site were not accessible by the OGI crew. At site 2, the percentage of OGI inaccessible emissions is around 9%-15%, lower than that in site 1.

Second, OGI is unable to detect all emission sources at a facility compared to aerial survey and has also been demonstrated across other supply chain segments.³⁹ In this study, when considering only those sources that OGI attempted to find, up to 50% or more of emissions identified by Bridger were not found by OGI at both sites.

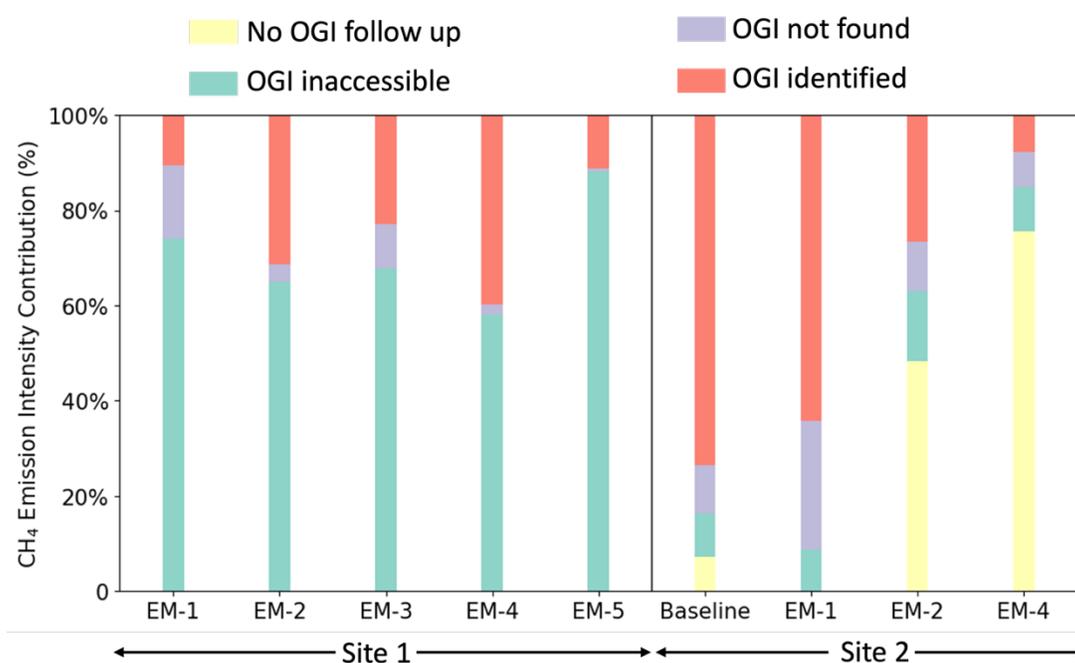


Figure 3. Emission reconciliation between ground based OGI follow-up survey and aerial Bridger survey. Confirming Bridger detections using OGI follow-up presents three options: (1) OGI identified possible sources (orange), (2) OGI did not find emissions found by Bridger (purple), (3) OGI could not access the source (green). Emissions found by Bridger that were not

followed up by OGI are shown in yellow. The y-axis represents the methane emission intensity contribution of each scenario of OGI/Bridger detection.

Table 1 shows the comparison of CH₄, CO₂ and total GHG emission intensity between GHGRP reporting and MII estimates at both sites. First, the 2022 GHGRP reported methane emission intensity is 74%-88% lower than time-averaged MII methane estimates for both sites. This observation demonstrates the underestimation of methane emissions in activity-based inventories, which is consistent with recent measurement studies across upstream and midstream segments of LNG supply chains. We also find that time-averaged MII relying solely on Bridger measurement are similar to time-averaged MII relying both on Bridger and ChampionX measurements, which indicates the consistence of methane emission estimates between Bridger and ChampionX measurements. In addition, we find that the GHGRP reported CO₂ emissions is consistent with time averaged MII CO₂ emission estimates for both sites. More importantly, since CO₂ emissions contribute to more than 95% of total GHG emissions (GWP-100 basis) for both sites, the GHGRP 2022 reported total GHG reasonably match time-averaged MII estimated GHG emissions to within measurement uncertainty. CO₂ EI at Site 1 is based on the OEI as measurements by ChampionX did not pass quality checks during analysis. Measured CO₂ EI at Site 2 is also consistent with satellite derived EIs at liquefaction terminals.⁴⁰

The range of methane EIs measured in this study are comparable to estimates from a recent empirical measurement campaign.³⁶ Comparisons with other non-empirical estimates of methane EI at liquefaction terminals are discussed in SI (SI section S10).

Table 1. Comparison of CH₄, CO₂ and total GHG emission intensity (GWP-100 basis) between GHGRP 2022 reporting and MII estimates at both sites. The values in the parentheses represent 95% confidence interval.

Site	GHG type	GHGRP 2022	Time-averaged MII (Bridger + Champion X)	Time-averaged MII (only Bridger based)
Site 1	CH ₄ (%)	0.005	0.019 (0.018-0.023)	0.019 (0.018-0.022)
	CO ₂ (kg CO ₂ /kg CH ₄ production)	0.217	0.232*	/
	CO₂e (kg CO₂e/kg CH₄ production)	0.218	0.238	/
Site 2	CH ₄ (%)	0.003	0.024 (0.022-0.028)	0.023 (0.022-0.028)
	CO ₂ (kg CO ₂ /kg CH ₄ production)	0.227	0.258 (0.218-0.299)	/
	CO₂e (kg CO₂e/kg CH₄ production)	0.228	0.265 (0.225-0.307)	/

*OEI used for CO₂ emissions estimates at Site 1 as direct measurements during baseline was rejected during QA/QC

Variability in methane EI at liquefaction terminals is a key feature of our measurement campaign. Furthermore, the complex three-dimensional nature of a liquefaction terminal where each liquefaction train could have over five levels makes ground based LDAR surveys time-consuming and ineffective. Safety and logistical considerations prevent ground crew from accessing several parts of the facility. Thus, aerial surveys that have been demonstrated to effectively estimate all emission sources above their detection threshold followed by ground based OGI to identify emitting component and initiate follow-up action can be an effective LDAR program. Even with aerial surveys, the complex nature of a liquefaction terminal will affect plume development. For example, a persistent fugitive emission source on the lower levels of a liquefaction train could appear intermittent on an aerial survey if the plume rise is blocked by higher levels of the train. Thus, interpreting aerial survey data requires an understanding of the underlying processes and operational information that can indicate the temporal nature of a

source. In this work, we took the conservative option to assume all sources are persistent – future work should focus on evaluating intermittency at these facilities.

This work provides the first measurement informed GHG EI estimates at US liquefaction terminals by employing multiscale measurement technologies. In addition, we demonstrate the role for aerial directed measurements in potentially informing operational maintenance for methane emission mitigation. Finally, we conclusively demonstrate that reported GHG EI of liquefaction terminals reasonably match measurement-informed GHG EI. This conclusion differs from every other segment of the natural gas supply chain, where MIIs are higher than reported inventory estimates. This is because carbon dioxide emissions, which can be accurately estimated using fuel emissions factors, comprise over 95% of all GHG emissions. Thus, in the context of developing LNG supply chain emissions intensity, the use of GHGRP-reported EI provides reasonably accurate estimates of total GHG emissions at liquefaction terminals.

ASSOCIATED CONTENT

Supporting Information

The following files are available free of charge.

Additional information referenced in the manuscript related to measurement technologies, measurement campaign, methane emissions intensity, Bridger measurement data analysis methodology, Bridger quantification uncertainty analysis, reconciliation between two aerial measurements and OEI, reconciliation between OEI and MII, flare measurements, GHG emission intensities under 20-year global warming potential, and GHG emission intensities normalized to LNG production (PDF).

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Author Contributions

F.C.G. conceived the study. A.P.R., G.R., S.R., and F.C.G. designed the measurement campaign. Y.Z. analyzed the field data, developed new models for measurement-informed inventories and uncertainty estimates, and wrote the paper. All authors contributed to the discussion and interpretation of results and reviewing the paper.

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Notes

The authors declare the following competing financial interest(s): F.G. and G.R. are currently employees of Cheniere Energy Inc. S.A.R.W was an employee of Cheniere till May 2024, and is currently an employee of SLR International. SLR International performs work for Cheniere, other oil and gas industry clients, academic institutions, and industry research organizations.

A.P.R. is currently a member of the Gas Pipeline Advisory Committee of the US Department of Transportation; in this role, he is a Special Government Employee. A.P.R. has current research support from the US Department of Energy, Environmental Defense Fund, and sponsors of the Energy Emissions Modeling and Data Lab (EEMDL).

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