

FLUXNET-CH₄ SYNTHESIS ACTIVITY

Objectives, Observations, and Future Directions

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We describe a new coordination activity and initial results for
a global synthesis of eddy covariance CH₄ flux measurements.

Atmospheric methane (CH₄) is the second-most important anthropogenic greenhouse gas following carbon dioxide (CO₂) (Myhre et al. 2013). The concentration of CH₄ in the atmosphere today is about 2.5 times higher than in 1750 (Saunois et al. 2016a). The increase in atmospheric CH₄ has arisen from human activities in agriculture, energy production, and waste disposal, and from changes in natural CH₄ sources and sinks (Saunois et al. 2016a,b, 2017; Turner et al. 2019). Based on top-down atmospheric inversions, global CH₄ emissions for the decade of 2003–12 were an estimated ~420 Tg C yr⁻¹ (range 405–426 Tg C yr⁻¹) (Saunois et al. 2016a). However, some analyses suggest that uncertainties in global CH₄ sources and sinks are higher than those for CO₂,

and uncertainties from natural sources exceed those from anthropogenic emissions (Saunois et al. 2016a). In particular, the largest source of uncertainty in the global CH₄ budget is related to emissions from wetlands and inland waters (Saunois et al. 2016a; Melton et al. 2013; Bastviken et al. 2011). Wetland CH₄ emissions may contribute as much as 25%–40% of the global total and are a leading source of interannual variability in total atmospheric CH₄ concentrations (Bousquet et al. 2006; Chen and Prinn 2006; Saunois et al. 2016a).

Direct, ground-based measurements of in situ CH₄ fluxes with high measurement frequency are important for understanding the responses of CH₄ fluxes to environmental factors including climate,

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for providing validation datasets for the land surface models used to infer global CH₄ budgets, and for constraining CH₄ budgets. Eddy covariance (EC) flux towers measure real-time exchange of gases such as CO₂, CH₄, water vapor, and energy between the land surface and the atmosphere. The EC technique has emerged as a widespread means of measuring trace gas exchange because it provides direct and near-continuous ecosystem-scale flux measurements without disturbing the soil or vegetation (Baldocchi 2003; Aubinet et al. 2012). There are more than 900 reported active and historical flux tower sites globally and approximately 7,000 site years of data collected (Chu et al. 2017). While most of these sites measure CO₂, water vapor, and energy exchange, the development of new and robust CH₄ sensors has resulted in a rapidly growing number of CH₄ EC measurements (Baldocchi 2014; Morin 2018), primarily in natural and agricultural wetlands (Petrescu et al. 2015).

Since the late 1990s, with a growing number of long-term, near-continuous EC measurements, the EC community has been well coordinated for integrating and synthesizing CO₂, water vapor and energy fluxes. This cross-site coordination resulted in the development of regional flux networks for Europe [EuroFlux, CarboEurope, and Integrated Carbon Observing System (ICOS)], Australia (OzFlux), North and South America (AmeriFlux, Large Biosphere Amazon, Fluxnet-Canada/Canadian Carbon Program, and MexFlux), Asia [AsiaFlux, ChinaFlux, KoFlux, and U.S.–China Carbon Consortium (USCCC)], and globally, FLUXNET (Papale et al. 2012; Baldocchi 2014). The resulting FLUXNET database (<http://fluxnet.fluxdata.org/>) has been used extensively to evaluate satellite measurements, inform Earth system models, generate data-driven CO₂ flux products, and provide answers to a broad range of questions about atmospheric fluxes related to ecosystems, land use and climate (Pastorello et al. 2017). FLUXNET has grown steadily over the past 25 years, enhancing our understanding of carbon, water and energy cycles in terrestrial ecosystems (Chu et al. 2017).

Similar community efforts and syntheses for CH₄ remain limited in part because EC measurements for CH₄ fluxes were rarer until recently. Whereas the earliest EC measurements of CO₂ fluxes date back to the late 1970s and early 1980s (Desjardins 1974; Anderson et al. 1984), the first EC CH₄ flux measurements only began in the 1990s (Verma et al. 1992; Shurpali and Verma 1998; Fan et al. 1992; Kim et al. 1999), with reliable, easy-to-deploy field sensors only becoming available in the past decade or so. EC CH₄ flux measurements became more feasible with advances

in sensor development, such as tunable diode laser absorption spectrometers, that allowed researchers to measure previously undetectable trace gas fluxes with higher signal to noise ratios (Rinne et al. 2007; McDermitt et al. 2011). After these new sensors were commercialized, and low-power, low-maintenance open-path sensors were developed that could be operated by solar panels in remote locations, the number of CH₄ flux tower measurements increased substantially (Baldocchi 2014; Morin 2018). The rapidly growing number of EC CH₄ flux measurements presents new opportunities for FLUXNET-type analyses and syntheses of ecosystem-scale CH₄ flux observations.

This manuscript describes initial results from a new coordination activity for flux tower CH₄ measurements organized by the Global Carbon Project (GCP) in collaboration with regional flux networks and FLUXNET. The goal of the activity is to develop a global database for EC CH₄ observations to answer regional and global questions related to CH₄ cycling. Here, we describe the objectives of the FLUXNET-CH₄ activity, provide an overview of the current geographic and temporal coverage of CH₄ flux measurements globally, present initial analyses exploring time scales of variability, uncertainty, trends, and drivers of CH₄ fluxes across 60 sites, and discuss future research opportunities for examining controls on CH₄ emissions and reducing uncertainties in the role of wetlands in the global CH₄ cycle.

FLUXNET-CH₄ SYNTHESIS OBJECTIVES AND TASKS. This activity is part of a larger GCP effort to establish and better constrain the global methane budget (www.globalcarbonproject.org/methanebudget/index.htm), and is designed to develop a CH₄ database component in FLUXNET for a global synthesis of CH₄ flux tower data. To this end, we are surveying, assembling, and synthesizing data from the EC community, in coordination with regional networks, including AmeriFlux's 2019 "Year of Methane" (<http://ameriflux.lbl.gov/year-of-methane/year-of-methane/>), FLUXNET initiatives, and other complementary activities. In particular, this work is being carried out in parallel with the EU's Readiness of ICOS for Necessities of Integrated Global Observations (RINGO) project, which is working to standardize protocols for flux calculations, quality control and gap-filling for CH₄ fluxes (Nemitz et al. 2018). Methane-specific protocols are needed because of the added complexities and high variability of CH₄ flux measurements and dynamics (Nemitz et al. 2018).

Our approach is to include all currently available and future CH₄ flux tower observations in a global CH₄ database, including freshwater, coastal, natural, and managed ecosystems, as well as upland ecosystems that may be measuring CH₄ uptake by soils. The initiative is open to all members of the EC community. Database compilation began in 2017 and is ongoing. Data from sites in the Americas can be submitted to AmeriFlux (<http://ameriflux.lbl.gov/data/how-to-uploaddownload-data/>); otherwise, data can be submitted to the European Fluxes Database Cluster (www.europe-fluxdata.eu/home/sites-list).

In addition to many applications, an ultimate goal of the FLUXNET-CH₄ activity is to generate a publicly available, open-access, data-driven global CH₄ emissions product using similar machine-learning-based approaches used for CO₂ fluxes (Jung et al. 2009; Tramontana et al. 2016). The product will be based on mechanistic factors associated with CH₄ emissions and new spatiotemporal information on wetland area and dynamics for constraining CH₄-producing areas. This gridded product will provide an independent bottom-up estimate of global wetland CH₄ emissions to compare with estimates of global CH₄ emissions from land surface models and atmospheric inversions. Recent work has shown the potential to upscale EC CH₄ flux observations across northern wetlands, with predictive performance comparable to previous studies upscaling net CO₂ exchange (Peltola et al. 2019); however, our focus is on a globally gridded product.

The near-continuous, high-frequency nature of EC measurements also offers significant promise for improving our understanding of ecosystem-scale CH₄ flux dynamics. As such, this synthesis also aims to investigate the dominant controls on net ecosystem-scale CH₄ fluxes from hourly to interannual time scales across wetlands globally, and to characterize scale-emergent, nonlinear, and lagged processes of CH₄ exchange.

Methane is produced during decomposition under anaerobic or reducing conditions and is transported to the atmosphere via plant-mediated transport, ebullition, and diffusion (Bridgham et al. 2013). During transport, CH₄ can pass through unsaturated soil layers and be consumed or oxidized by aerobic bacteria (Wahlen 1993). Process-based biogeochemical models developed and applied at site, regional, and global scales simulate these individual processes with varying degrees of complexity (Bridgham et al. 2013; Melton et al. 2013; Poulter et al. 2017; Castro-Morales et al. 2018; Grant and Roulet 2002). The large range in predicted wetland CH₄ emissions rates suggests that there is both

substantial parameter and structural uncertainty in large-scale CH₄ flux models, even after accounting for uncertainties in wetland areas (Poulter et al. 2017; Saunois et al. 2016a; Melton et al. 2013; Riley et al. 2011). A global EC CH₄ database and associated environmental variables can help constrain the parameterization of process-based biogeochemistry models (Saunois et al. 2016a; Bridgham et al. 2013; Oikawa et al. 2017). Furthermore, a key challenge is evaluating globally applicable process-based CH₄ models at a spatial scale comparable to model grid cells (Melton et al. 2013; Riley et al. 2011). A globally gridded wetland CH₄ emissions product upscaled from EC fluxes can help resolve this issue by providing a scale-appropriate model evaluation dataset. As such, the global CH₄ database and gridded product will also be used to parameterize and benchmark the performance of land surface models of global CH₄ emissions, providing a unique opportunity for informing and validating biogeochemical models.

METHODS. Based on a survey of the EC community (announced via the fluxnet-community@george.lbl.gov and AmeriFlux-Community@lbl.gov listservs), information available in regional networks and FLUXNET, and the scientific literature, we estimate that at least 200 sites worldwide are currently applying the EC method for CH₄ flux measurements (Fig. 1). Here we focus on findings from across 60 of the ~110 sites currently committed to participating in our FLUXNET-CH₄ activity [Table A1 in the appendix and Table ES1 in the online supplemental material (<https://doi.org/10.1175/BAMS-D-18-0268.2>)]. Data from this initial set of sites were selected because they were publicly available or were contributed directly by site principal investigators (PIs). We will continue to engage the EC community more broadly and expand the database in the future.

Data standardization, gap-filling, and partitioning. We used similar data processing procedures as FLUXNET to standardize and gap-fill measurements, and in the case of net CO₂ exchange, partition fluxes across sites (<http://fluxnet.fluxdata.org/data/aboutdata/data-processing-101-pipeline-and-procedures/>). Standard quality assurance and quality control of the data were first performed by site PIs. In nearly all cases, data collected by the local tower teams were first submitted to the data archives hosted by the regional flux networks, where data are prescreened and formatted based on the regional network data protocols. Data from the regional networks then entered our flux processing procedure.

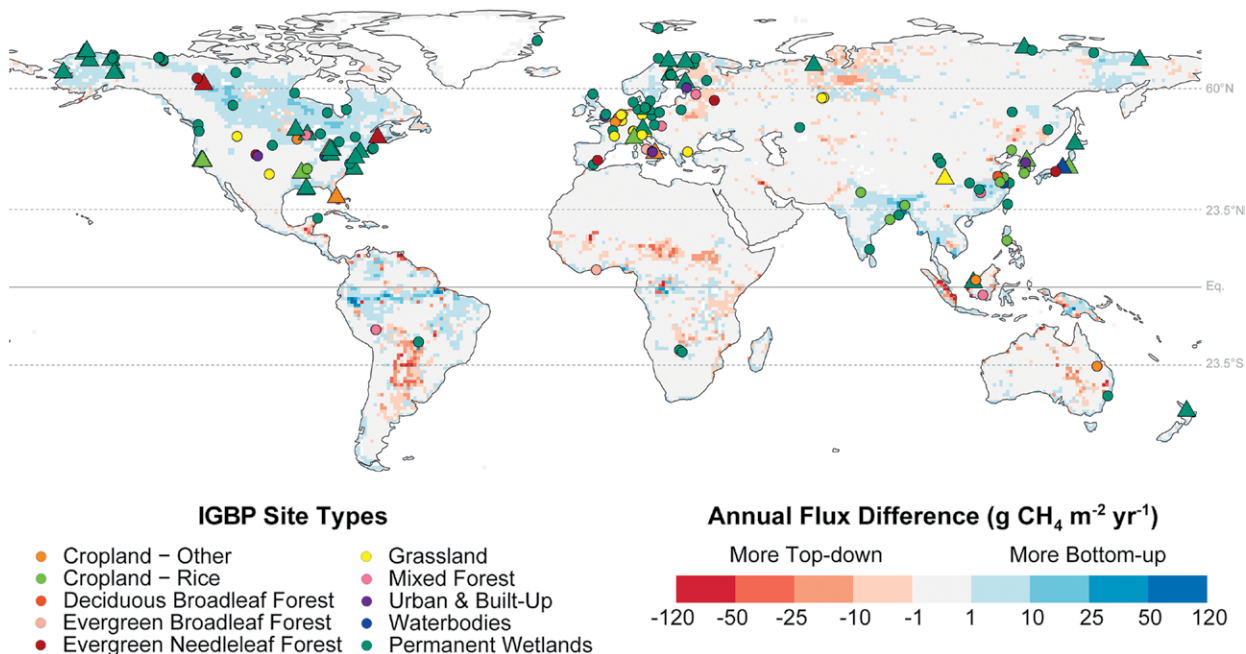


FIG. 1. Location of the 200 tower sites that report eddy covariance CH_4 flux measurements worldwide. Triangles indicate sites from which data are included in this manuscript, with circles indicating additional flux towers measuring CH_4 emissions. The colors of the markers represent the vegetation type based on the International Geosphere-Biosphere Programme (IGBP) definition. See Table ESI for a list of sites, their characteristics, and years of operation. Sites are overlaid over a map of the differences between the average CH_4 emissions over 2000–10 between top-down and bottom-up wetland CH_4 estimates. Top-down estimates are represented by the natural fluxes inventoried in NOAA's CarbonTracker (www.esrl.noaa.gov/gmd/ccgg/carbontracker-ch4/). Bottom-up emissions were produced from an ensemble of 11 Earth system model simulations (Poulter et al. 2017).

Within our processing procedure, data were first checked for obvious problems including unit errors, spikes, and out-of-range values based on visualization of the data and statistical metrics. Next, the data were filtered, gap-filled, and partitioned. Friction velocity (u_*) filtering, based on relating nighttime CO_2 fluxes to u_* , was implemented using the R EddyProc package (Wutzler et al. 2018) for R statistical software (R Core Team 2018, version 3.5.0), although in a few cases u_* filtering was performed by the site PIs. Gaps in meteorological variables including air temperature (TA), incoming shortwave (SW_{IN}) and longwave (LW_{IN}) radiation, vapor pressure deficit (VPD), pressure (PA), precipitation (P), and wind speed (WS) were filled with ERA-Interim (ERA-I) reanalysis data (Vuichard and Papale 2015). Gaps in CO_2 and latent and sensible heat fluxes were filled using the marginal distribution sampling method (Reichstein et al. 2005) using the R EddyProc package (Wutzler et al. 2018). Net CO_2 fluxes were partitioned into gross primary production (GPP) and ecosystem respiration (ER) using both the nighttime (Reichstein et al. 2005) and daytime (Lasslop et al. 2010) approaches also implemented in R EddyProc (Wutzler et al. 2018).

There are as yet no standards for gap-filling CH_4 flux measurements and this is an active and ongoing area of research (Nemitz et al. 2018). Gaps in CH_4 fluxes were filled using artificial neural networks (ANNs), as they have shown good performance for gap-filling CH_4 flux data (Dengel et al. 2013; Knox et al. 2015; Morin et al. 2014a; Nemitz et al. 2018; Goodrich et al. 2015). Details of the ANN routine are provided in Knox et al. (2016) and are summarized here briefly. The ANN routine was optimized for both generalizability and representativeness. To facilitate representativeness, explanatory data were divided into a maximum of 15 data clusters using the k -means algorithm. To avoid biasing toward conditions with better flux data coverage (e.g., summer and daytime), data used to train, test, and validate the ANN were proportionately sampled from these clusters. Several neural network architectures of increasing complexity were tested, ranging from one hidden layer with the number of nodes equal to the number of explanatory data variables (N) to two hidden layers with $1.5N$ and $0.75N$ nodes, respectively. The architecture of each neural network was initialized 10 times with random starting weights, and the initialization resulting in the lowest mean sampling

error was selected. The simplest architecture, whereby additional increases in complexity resulted in <5% reduction in mean squared error, was chosen and the prediction saved. This procedure was repeated with 20 resamplings of the data, and missing half hours were filled using the median prediction. A standard set of variables available across all sites were used to gap-fill CH₄ fluxes (Dengel et al. 2013), including TA, SW_{IN}, WS, PA, and sine and cosine functions to represent seasonality. These meteorological variables were selected since they are relevant to CH₄ exchange and were gap-filled using the ERA-I reanalysis data. Other variables related to CH₄ exchange such as water table depth (WTD) or soil temperature (TS) were not included as explanatory variables as they were not available across all sites or had large gaps that could not be filled using the ERA-I reanalysis data. These missing data for variables highlight some of the key challenges in standardizing CH₄ gap-filling methods across sites and emphasize the need for standardized protocols of auxiliary measurements across sites (cf. “Future research directions and needs” section) (Nemitz et al. 2018; Dengel et al. 2013). ANN gap-filling was performed using MATLAB (MathWorks 2018, version 9.4.0).

Annual CH₄ budgets represent gap-filled, half-hourly fluxes integrated over an entire year or growing season. If fluxes were only measured during the growing season, we assumed that fluxes outside of this period were negligible, although we acknowledge that cold season fluxes can account for as much as ~13%–50% of the annual CH₄ emissions in some locations (Zona et al. 2016; Treat et al. 2018b; Helbig et al. 2017a; Kittler et al. 2017).

Uncertainty estimation. ANNs were also used to estimate annual gap-filled and random uncertainty in CH₄ flux measurements (Richardson et al. 2008; Moffat et al. 2007; Anderson et al. 2016; Knox et al. 2018). Here, we focus on assessing the random error, but a full assessment of total flux measurement error also requires quantifying systematic error or bias (Baldocchi 2003). Systematic errors, due to incomplete spectral response, lack of nocturnal mixing, submesoscale circulations, and other factors are discussed elsewhere (Baldocchi 2003; Peltola et al. 2015) and are the focus of other ongoing initiatives.

Random errors in EC fluxes follow a double exponential (Laplace) distribution with a standard deviation varying with flux magnitude (Richardson et al. 2012, 2006). Model residuals of gap-filling algorithms such as ANNs provide a reliable, and conservative “upper limit,” estimate of the random

flux uncertainty (Moffat et al. 2007; Richardson et al. 2008). For half-hourly CH₄ flux measurements, random error was estimated using the residuals of the median ANN predictions. At each site, the probability density function (PDF) of the random flux measurement error more closely followed a double-exponential (Laplace) rather than normal (Gaussian) distribution, with the root-mean-square error (RMSE) for the Laplace distribution fitted to the PDF of random errors consistently lower than the normal distributed error. From half-hourly flux measurements, random error can also be estimated using the daily differencing approach (Richardson et al. 2012). Random error estimates [$\sigma(\delta)$], as expressed as the standard deviation of the double-exponential distribution with scaling parameter β , where $\sigma(\delta) = \sqrt{2}\beta$ (Richardson et al. 2006), were found to be nearly identical using the two approaches [$\sigma(\delta)_{\text{model_residual}} = 1.0 \times \sigma(\delta)_{\text{daily_differencing}} + 1.21$; $r^2 = 0.97$, $p < 0.001$], supporting the use of the model residual approach for estimating random error. As discussed below, $\sigma(\delta)$ scaled linearly with the magnitude of CH₄ fluxes at nearly all sites. To quantify random uncertainty of cumulative fluxes, we used a Monte Carlo simulation that randomly draws 1,000 random errors for every original measurement using $\sigma(\delta)$ binned by flux magnitude, and then computed the variance of the cumulative sums (Anderson et al. 2016). For gap-filled values, the combined gap-filling and random uncertainty was calculated from the variance of the cumulative sums of the 20 ANN predictions (Anderson et al. 2016; Oikawa et al. 2017; Knox et al. 2015). The annual cumulative uncertainty at 95% confidence was estimated by adding the cumulative gap-filling and random measurement uncertainties in quadrature (Richardson and Hollinger 2007; Anderson et al. 2016). Note that when reporting mean or median annual CH₄ fluxes across sites, error bars represent the standard error.

Wavelet-based time-scale decomposition. Methane fluxes are highly dynamic and vary across a range of time scales (Sturtevant et al. 2016; Koebisch et al. 2015). For example, in wetlands with permanent inundation, the seasonal variation of CH₄ exchange is predominantly controlled by temperature and plant phenology (Chu et al. 2014; Sturtevant et al. 2016). Ecosystem CH₄ exchange also varies considerably at both longer (e.g., interannual; Knox et al. 2016; Rinne et al. 2018) and shorter (e.g., weeks, days, or hours; Koebisch et al. 2015; Hatala et al. 2012; Schaller et al. 2018) time scales. Wavelet decomposition is a particularly useful tool for investigating scale in geophysical

and ecological analysis (Cazelles et al. 2008; Torrence and Compo 1998), because it can characterize both the time scale and location of patterns and perturbations in the data. Partitioning variability across temporal scales can help to isolate and characterize important processes (Schaller et al. 2018).

The maximal overlap discrete wavelet transform (MODWT) was used to decompose the time scales of variability in gap-filled CH_4 flux measurements, as described in Sturtevant et al. (2016). The MODWT allows the time series to be decomposed into the detail added from progressively coarser to finer scales and either summed or treated individually to investigate patterns across scales. We reconstructed the detail in the fluxes for dyadic scales 1 (2^1 measurements = 1 h) to 14 (2^{14} measurements = 341 days). Since patterns generated by ecological processes tend to occur over a scale range rather than at one individual scale, the detail over adjacent scales were summed to analyze four general time scales of variation (Sturtevant et al. 2016). These time scales included the “hourly” scale (1–2 h) representing perturbations such the passage of clouds overhead and turbulent scales up to the spectral gap, the “diel” scale (4 h to 1.3 days) encompassing the diel cycles in sunlight and temperature, the “multiday” scale (2.7 to 21.3 days) reflecting synoptic weather variability or fluctuations in water levels, and the “seasonal” scale (42.7 to 341 days) representing the annual solar cycle and phenology. Data were wavelet decomposed into the hourly, diel, and multiday scales using the Wavelet Methods for Time Series Analysis (WMTSA) Wavelet Toolkit in MATLAB.

Statistical analysis. We tested for significant relationships between log-transformed annual CH_4 emissions and a number of covariates using linear mixed-effects models as described in Treat et al. (2018b). The predictor variables of CH_4 flux we evaluated included: biome or ecosystem type (categorical variables), and continuous biophysical variables including mean seasonal WTD, mean annual soil and air temperature (T_{MST} and T_{MAT} , respectively), net ecosystem exchange (NEE), GPP, and ER. When considering continuous variables, we focused on freshwater wetlands for comparison with previous CH_4 synthesis activities. Soil temperature was measured between 2 and 25 cm below the surface in different studies. The results below are presented for GPP and ER covariates that are partitioned using the nighttime flux partitioning algorithm (Wutzler et al. 2018; Reichstein et al. 2005), although similar findings were obtained using daytime partitioned estimates. Additionally, individual sites or

site years were excluded when gaps in measurements exceeded two consecutive months, which explains the differences in the number of sites and site years in the “Environmental controls on annual CH_4 emissions across freshwater wetland sites” section below.

Mixed-effects modeling was used because of the potential bias of having measurements over several years, with site included as a random effect in the analysis (Treat et al. 2018b). The significance of individual predictor variables was evaluated using a χ^2 test against a null model using only site as a random variable (Bates et al. 2015), with both models fit without reduced maximum likelihood. For multiple linear regression models, we used the model selection process outlined in Zuur et al. (2009). To incorporate annual cumulative uncertainty when assessing the significance of trends and differences in annual CH_4 fluxes across biomes and ecosystem types, we used a Monte Carlo simulation that randomly draws 1,000 annual cumulative uncertainties for each estimate of annual CH_4 flux. For each random draw the significance of the categorical variable was tested using a χ^2 test against the null model with only site as a random variable. We report the marginal r^2 (r_m^2), which describes the proportion of variance explained by the fixed factors alone (Nakagawa and Schielzeth 2013). The mixed-effects modeling was implemented using the lmer command from the lme4 package (Bates et al. 2015) for R statistical software.

RESULTS AND DISCUSSION. *Geographic and temporal coverage of eddy covariance CH_4 flux measurements.* We identified 200 sites worldwide that are applying the EC method for CH_4 (Fig. 1; Table ES1); wetlands (including natural, managed, and restored wetlands) comprise the majority of sites (59%), with rice agriculture (10%) as the second-most represented vegetation type. The predominance of wetland and rice paddy sites in the database is unsurprising because many studies are designed to target ecosystems expected to have relatively large CH_4 emissions. However, there are also sites in ecosystems that are typically smaller sources or even sinks of CH_4 such as upland forests (13%) and grasslands (8%). Additionally, six sites (~3%) are urban, with another five sites measuring CH_4 fluxes from open water bodies. Although identified sites span all continents except Antarctica, the majority are concentrated in North America and Europe, with a growing number of sites in Asia (Fig. 1; Table ES1).

Measurements of CH_4 fluxes cover a broad range of climates and a large fraction of wetland habitats (Fig. 2), with the tropics and tropical wetlands notably

underrepresented. As discussed below (see “Future research directions and needs” section), one important goal of FLUXNET and the regional networks is to increase site representativeness and extend measurements in undersampled regions. Increasing the number of tropical sites is particularly important for CH₄ because more than half of global CH₄ emissions are thought to come from this region (Saunois et al. 2016a; Dean et al. 2018). Furthermore, compared to northern wetlands, their biogeochemistry remains relatively poorly understood (Mitsch et al. 2009; Pangala et al. 2017). We expect the number of CH₄ flux sites and their geographic and temporal coverage to continue to increase, as has occurred through time for CO₂, water vapor, and energy flux measurements in FLUXNET (Pastorello et al. 2017; Chu et al. 2017).

Long-term CH₄ flux time series are key to understanding the causes of year-to-year variability and

trends in fluxes (Chu et al. 2017; Euskirchen et al. 2017; Pugh et al. 2018). The longest continuous record of CH₄ flux measurements, from a fen in Finland (Rinne et al. 2018), is now ~14 years and ongoing (Table ES1). Three other sites have measurements exceeding 10 years; however, the median length is 5 years, with most sites established from 2013 onward (Table ES1). Longer time series are also important for both exploring the short- and long-term effects of extreme events on fluxes and tracking the response of disturbed or restored ecosystems over time (Pastorello et al. 2017). Furthermore, they can help address new and emerging science questions, such as quantifying CH₄ feedbacks to climate with rising temperatures and associated changes in ecosystem composition, structure and function (Helbig et al. 2017a,b; Dean et al. 2018), and the role of wetland emissions in atmospheric CH₄ variability (McNorton et al. 2016; Poulter et al. 2017).

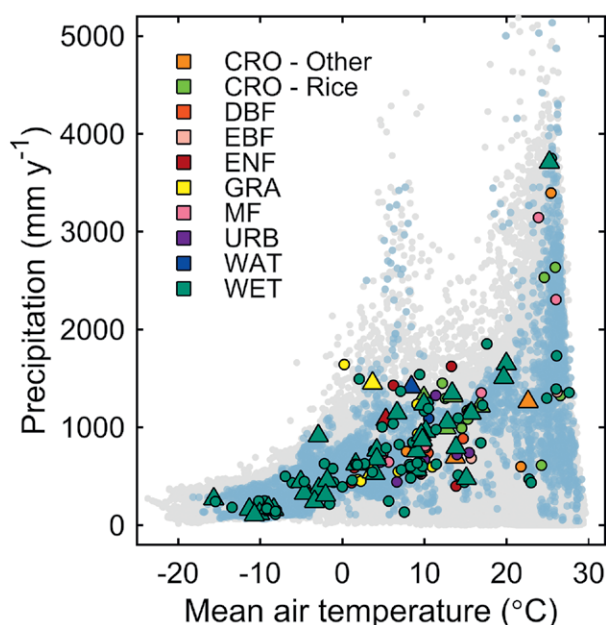


FIG. 2. Distribution of sites by mean annual air temperature and precipitation. Tower locations are shown as circles or triangles (see Fig. 1), with vegetation type in color based on the IGBP definitions (CRO = croplands; DBF = deciduous broadleaf forests; EBF = evergreen broadleaf forests; ENF = evergreen needleleaf forests; GRA = grasslands; MF = mixed forests; URB = urban and built-up lands; WAT = water bodies; WET = permanent wetlands). Gray dots represent annual mean temperature and total precipitation from the CRU TS 3.10 gridded climate dataset over the entire landmass (Harris et al. 2014), whereas blue dots represent grid cells with >25% wetland fraction as estimated using the Global Lakes and Wetlands Database (Lehner and Döll 2004). Temperature and precipitation grid cells included in this figure were averaged from 1981 to 2011, at 0.5° resolution.

CH₄ fluxes and trends across biomes and ecosystem types. Half-hourly and annual net CH₄ fluxes for the 60 sites currently included in the database exhibited strong variability across sites (Figs. 3 and 4). Across the dataset, the mean half-hourly CH₄ flux was greater than the median flux, indicating a positively skewed distribution with infrequent, large emissions (Fig. 3a), similar to findings from chamber-based syntheses (Olefeldt et al. 2013; Turetsky et al. 2014). Mean and median CH₄ fluxes were smaller at higher latitudes and larger at lower latitudes (Fig. 3b), comparable again to trends in CH₄ fluxes observed in predominantly chamber-based syntheses (Bartlett and Harriss 1993; Turetsky et al. 2014; Treat et al. 2018b).

The continuous nature of EC flux measurements is well suited for quantifying annual ecosystem-scale CH₄ budgets, along with accumulated uncertainty (cf. “Gap-filling performance and uncertainty quantification” section). Annual estimates of net CH₄ flux for each of the 60 sites in the flux tower database ranged from $-0.2 \pm 0.02 \text{ g C m}^{-2} \text{ yr}^{-1}$ for an upland forest site to $114.9 \pm 13.4 \text{ g C m}^{-2} \text{ yr}^{-1}$ for an estuarine freshwater marsh (Rey-Sanchez et al. 2018), with fluxes exceeding $40 \text{ g C m}^{-2} \text{ yr}^{-1}$ at multiple sites (Fig. 4b). These emissions are of a considerably broader range and have much higher annual values than in an earlier synthesis by Baldocchi (2014), which included published values from 13 sites (Fig. 4a); median annual CH₄ fluxes ($\pm$ SE) in that study were $6.4 \pm 1.9 \text{ g C m}^{-2} \text{ yr}^{-1}$, compared with $10.0 \pm 1.6 \text{ g C m}^{-2} \text{ yr}^{-1}$ for our expanded database. Annual CH₄ sums in our database were positively skewed, with skewness increasing with additional observations due largely to the inclusion of

high CH_4 -emitting freshwater marsh sites (Fig. 4).

As suggested from Fig. 3b, annual wetland CH_4 emissions differed significantly among biomes, even when considering accumulated uncertainty [average Monte Carlo $\chi^2 = 13.4$ (12.1–14.7, 95% confidence interval), degrees of freedom (df) = 3, $p < 0.05$] (Table 1). Median CH_4 emissions were significantly lower for tundra wetlands ($2.9 \pm 1.3 \text{ g C m}^{-2} \text{ yr}^{-1}$) than temperate wetlands ($27.4 \pm 3.4 \text{ g C m}^{-2} \text{ yr}^{-1}$). Higher CH_4 emissions were observed from subtropical/tropical wetlands ($43.2 \pm 11.2 \text{ g C m}^{-2} \text{ yr}^{-1}$), based on only three site years of data; however, emphasizing the need for additional flux tower measurements in the tropics.

Whereas annual boreal/taiga wetland CH_4 emissions were comparable to values reported in a recent synthesis of predominantly chamber-based CH_4 flux measurements (Treat et al. 2018b), our tower-based measurements are ~50% lower and over 6 times higher for tundra and temperate wetlands, respectively (Table 1). The inconsistencies highlighted in Table 1 not only reflect the differences in the number and location of sites between datasets, but also the discrepancies resulting from different measurement

techniques. Several studies have noted considerable differences in CH_4 emissions measured using EC and chamber techniques, with estimates from chambers often higher than those from the EC measurements (Schrier-Uijl et al. 2010; Hendriks et al. 2010; Meijide et al. 2011; Krauss et al. 2016). This distinction highlights the need for additional studies investigating the systematic differences caused by the different spatial and temporal sampling footprints of these methods (Krauss et al. 2016; Morin et al. 2017;

Windham-Myers et al. 2018; Xu et al. 2017). Characterizing discrepancies between measurement techniques may also help constrain bottom-up estimates of CH_4 emissions and reduce the disagreement of ~15 Tg C yr^{-1} between bottom-up (139 Tg $\text{CH}_4 \text{ yr}^{-1}$) and top-down (125 Tg $\text{CH}_4 \text{ yr}^{-1}$) estimates of CH_4 emissions from natural wetlands (Saunois et al. 2016a).

Annual CH_4 emissions also differed significantly across ecosystems [average Monte Carlo $\chi^2 = 45.5$ (39.3–50.1), df = 9, $p < 0.001$; Fig. 5], with median fluxes highest for freshwater marshes ($43.2 \pm$

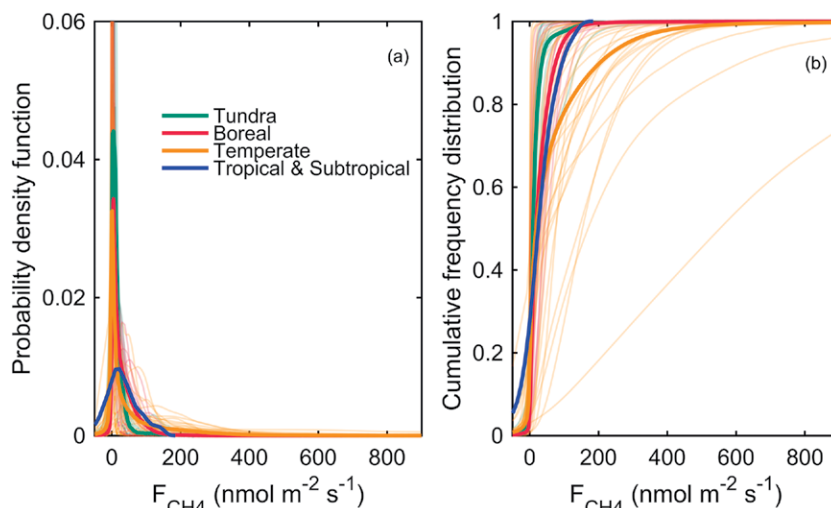


FIG. 3. (a) Probability density function, and (b) cumulative frequency distribution of half-hourly CH_4 flux (F_{CH_4}) data for sites currently included in the database (60 sites) aggregated by biome. Thin lines represent individual sites, whereas thicker lines present sites aggregated by biome. All cases are approximated by kernel density estimation. Note that whereas the x axis is scaled between -50 and $900 \text{ nmol m}^{-2} \text{ s}^{-1}$ for visualization purposes, some CH_4 fluxes exceed this range.

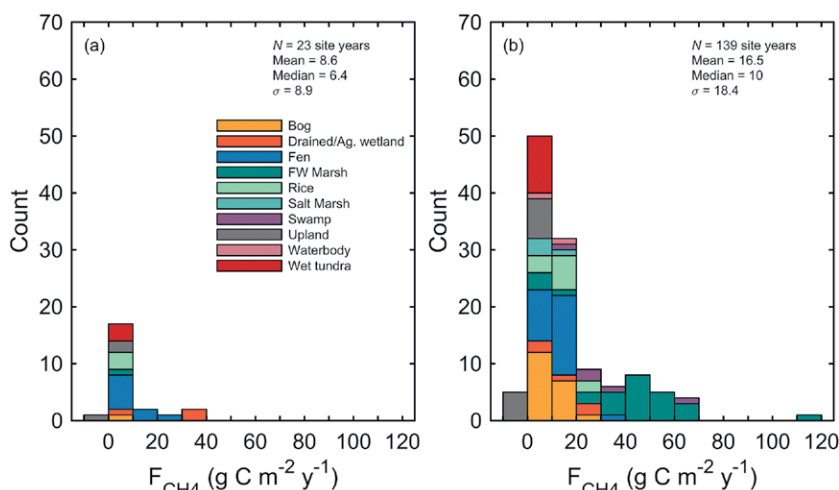


FIG. 4. (a) Histogram of annual CH_4 fluxes (F_{CH_4} ; $\text{g C m}^{-2} \text{ yr}^{-1}$) measured with eddy covariance and published in the synthesis by Baldocchi (2014), and (b) histogram of our annual CH_4 fluxes including additional site years of data estimated from the 60 sites listed in Table A1.

TABLE 1. Number of site years and characteristics of CH₄ fluxes (g C m⁻² yr⁻¹) currently included in the database. Fluxes are compared with measurements reported in a recent synthesis of predominantly chamber-based CH₄ flux measurements. Biome type was extracted from Olson et al. (2001) using site coordinates and includes tundra, boreal/taiga, temperate, and tropical/subtropical. Wetland CH₄ emissions differed significantly across biomes, with letters indicating significant differences ($\alpha = 0.05$) among biomes. Note that similar to our tower only dataset, values from Treat et al. (2018b) represent measured annual fluxes derived from a smaller dataset where measurements were made in the growing season and nongrowing season.

Biome	No. of site years	Median annual CH ₄ flux	25th percentile	75th percentile	References
Tundra	10	2.9	1.8	4.2	This study—All sites
	10	2.9 ^a	1.8	4.2	This study—Wetlands
	31	5.6	1.0	11.4	Treat et al. (2018b)—All sites
	26	6.3	3.0	16.4	Treat et al. (2018b)—Wetlands
Boreal and taiga	35	8.3	4.1	10.9	This study—All sites
	30	9.5 ^{ab}	6.0	11.3	This study—Wetlands
	68	13.1	3.5	23.7	Treat et al. (2018b)—All sites
	67	13.2	3.6	23.7	Treat et al. (2018b)—Wetlands
Temperate	72	16.4	7.9	35.9	This study—All sites
	47	27.4 ^b	10.0	47.3	This study—Wetlands
	27	4.3	0.3	41.7	Treat et al. (2018b)—All sites
	25	5.3	0.8	42.2	Treat et al. (2018b)—Wetlands
Tropical and subtropical	3	43.2	20.0	46.8	This study—All sites
	3	43.2 ^{ab}	20.0	46.8	This study—Wetlands
	—	—	—	—	Treat et al. (2018b)—All sites
	—	—	—	—	Treat et al. (2018b)—Wetlands

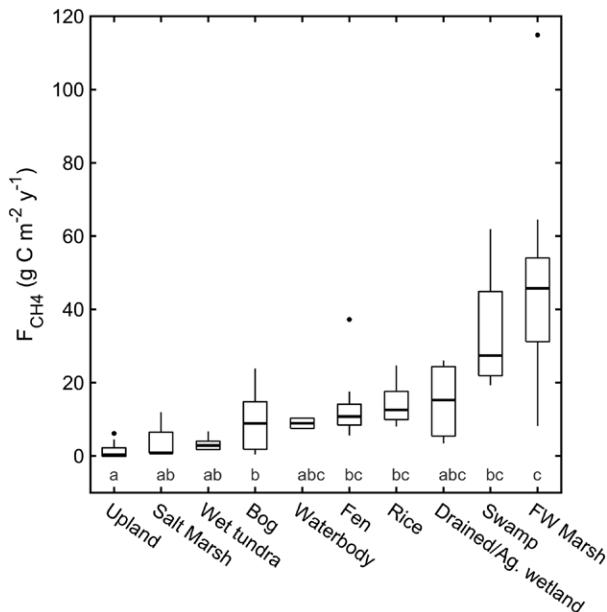


FIG. 5. Annual CH₄ fluxes (F_{CH_4} ; g C m⁻² yr⁻¹) among ecosystem types for the 60 sites currently included in the database (Table A1). Letters indicate significant differences ($\alpha = 0.05$) among ecosystem types. Median value, first quartile, and third quartile are presented in the boxes, and dots represent outliers, which are defined as observations more than 1.5 times the interquartile range away from the top or bottom of the box.

4.2 g C m⁻² yr⁻¹) and lowest for upland ecosystems (1.3 ± 0.7 g C m⁻² yr⁻¹). Treat et al. (2018b) also observed the highest annual emissions in marshes and reported a similar median value for temperate marshes (49.6 g C m⁻² yr⁻¹). Wet tundra and bogs had significantly lower annual emissions than marshes (Fig. 5), which in part reflects their presence in colder boreal and tundra systems, as well as differences in vegetation type, nutrient status, and hydrological regime (Treat et al. 2018b). Low median CH₄ emission was observed from salt marshes in our dataset (0.8 ± 2.9 g C m⁻² yr⁻¹), because high sulfate concentrations inhibit methanogenesis (Poffenbarger et al. 2011; Holm et al. 2016). Even drained wetlands converted to agricultural land can be large sources of CH₄ associated with seasonal flooding (Fig. 5). Median annual CH₄ flux from rice was 12.6 ± 1.6 g C m⁻² yr⁻¹, which is slightly lower than the IPCC default value of 15 g C m⁻² yr⁻¹ (Sass 2003).

Environmental controls on annual CH₄ emissions across freshwater wetland sites. Using an integrated CH₄ flux database, we can begin to investigate the factors associated with varying CH₄ emissions across sites. We explored the effects of WTD, T_{MST} or T_{MAT} , NEE, GPP, and ER on annual CH₄ flux. At global scales,

T_{MAT} and T_{MST} were the most important predictors of annual CH_4 flux across wetland sites ($p < 0.001$ for each), with the fixed factor of T_{MAT} or T_{MST} explaining $\sim 65\%$ of the variation in log transformed annual CH_4 emission (Figs. 6a,b). Previous synthesis studies also observed a significant, but weaker, relationship between soil temperature and average CH_4 emissions across sites, explaining $<15\%$ of the variation in CH_4 flux in those studies (Olefeldt et al. 2013; Yvon-Durocher et al. 2014). However, our findings are consistent with numerous site-level studies that report a strong correlation between wetland CH_4 emissions and temperature, with nearly 95% of all EC studies reporting a significant relationship between temperature and CH_4 flux (Morin 2018). Across sites, Peltola et al. (2019) found that the most important predictor in a random forest model used to upscale EC CH_4 emissions across northern latitudes was temperature, again highlighting the importance of temperature in regulating CH_4 emissions within and across sites.

Water table depth has also commonly been identified as a key control on CH_4 emissions (Turetsky et al. 2014; Bubier et al. 2005), because higher water levels often inhibit oxygen availability and lower the soil reduction potential, making methanogenesis more thermodynamically favorable. Although predominantly chamber-based wetland CH_4 syntheses have found a positive relationship between WTD and average or annual CH_4 emissions across sites (Olefeldt et al. 2013; Turetsky et al. 2014; Treat et al. 2018b), we observed no significant relationship between mean WTD and annual CH_4 flux across all sites ($\chi^2 = 0.2$, $df = 1$, $p = 0.66$, $N_{\text{sites}} = 20$, $N_{\text{site_yr}} = 46$), even when considering WTD^2 or WTD^3 (Olefeldt et al. 2013). However, if we consider only sites where WTD was below the soil surface for part or all of the year (Fig. 6c, solid circles), we did observe a significant relationship with WTD ($p < 0.05$). Conversely, CH_4 emissions for permanently inundated sites showed no

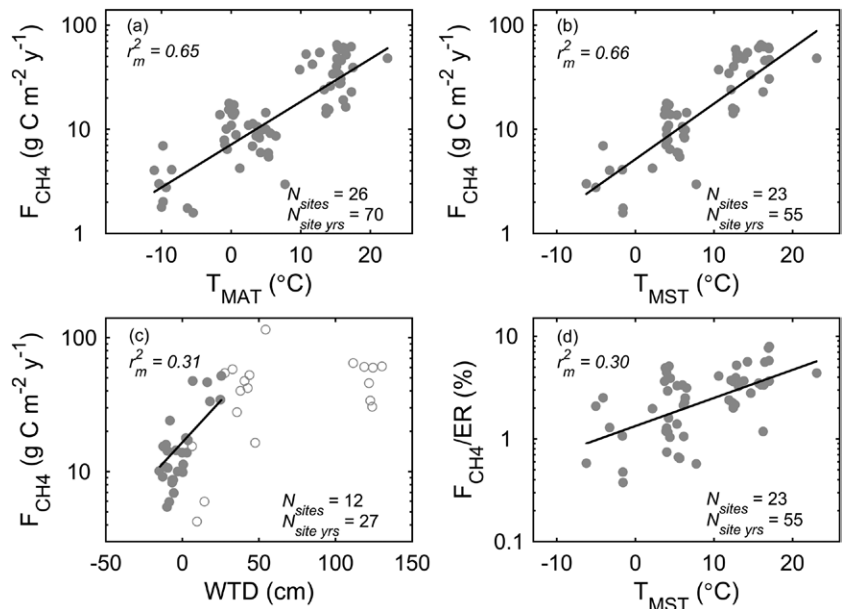


Fig. 6. Relationship between annual CH_4 flux (F_{CH_4}) and (a) mean annual air temperature (T_{MAT}) ($\chi^2 = 36.7$, $df = 1$, $p < 0.001$), (b) mean annual soil temperature (T_{MST}) ($\chi^2 = 32.3$, $df = 1$, $p < 0.001$) for freshwater wetlands, and (c) mean water table depth (WTD). While there was no significant relationship between mean annual WTD and annual CH_4 flux across all sites, there was a significant relationship if we consider only sites where WTD was below the soil surface for part or all of the year (solid circles) ($\chi^2 = 5.6$, $df = 1$, $p < 0.05$). Open circles in (c) indicate CH_4 emissions for permanently inundated sites. (d) Temperature dependence of the annual CH_4/ER ratio ($\chi^2 = 12.0$, $df = 1$, $p < 0.001$). Lines represent the fitted values for the population.

significant relationship with WTD (Fig. 6c, open circles) ($\chi^2 = 0.5$, $df = 1$, $p = 0.50$, $N_{\text{sites}} = 13$, $N_{\text{site_yr}} = 19$). This result supports the finding that wetlands that are permanently inundated or exhibit little variation in WTD tend to show weak to no correlation between WTD and CH_4 emissions (Chu et al. 2014; Jackowicz-Korczyński et al. 2010; Rinne et al. 2007; Christensen et al. 2003); in contrast, wetlands with lower and more variable water levels often have a significant relationship between WTD and CH_4 emissions (Bubier et al. 2005; Treat et al. 2007). However, only half of the sites currently included in the database report water table position, and given the importance of WTD in regulating CH_4 exchange, it is critical to ensure that WTD is measured across all sites.

Gross primary production and ER were both significant positive predictors of annual CH_4 flux ($\chi^2 = 21.3$, $df = 1$, $p < 0.001$, $r^2_m = 0.29$ and $\chi^2 = 17.1$, $df = 1$, $p < 0.001$, $r^2_m = 0.25$, respectively, $N_{\text{sites}} = 26$, $N_{\text{site_yr}} = 64$), although there was no significant relationship between NEE and annual CH_4 flux ($\chi^2 = 0.9$, $df = 1$, $p = 0.33$, $N_{\text{sites}} = 2$, $N_{\text{site_yr}} = 64$). However, when considering GPP or ER in a multiple linear regression model with T_{MST} , including interaction terms (Chu

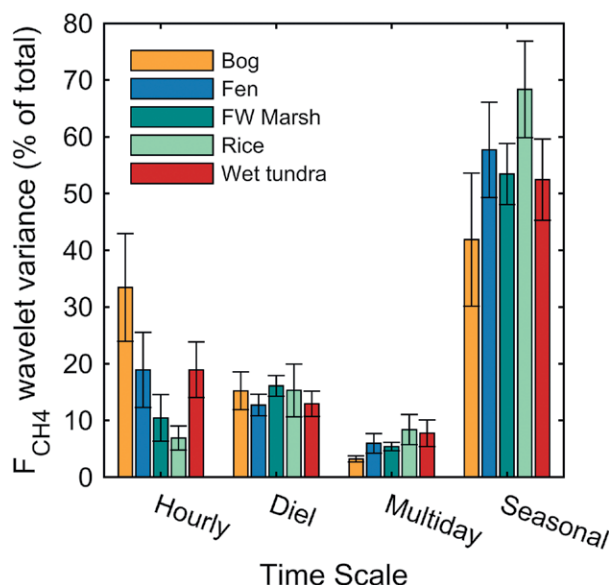


FIG. 7. Variance of CH₄ flux (F_{CH_4}) wavelet coefficients across time scales, as a percentage of the total variance, averaged by wetland type. Error bars represent the standard error. Note that only ecosystem types with at least 6 sites are shown here, including bogs, fens, freshwater (FW) marshes, rice paddies, and wet tundra.

et al. 2014), neither GPP nor ER were significant, suggesting that the observed relationship with GPP or ER was due to covariation with soil temperature and, possibly, other environmental drivers.

The strong temperature dependence of ecosystem-scale CH₄ emissions we observed across wetland sites is in line with the high temperature sensitivity of CH₄ emissions found across microbial to ecosystem scales (Yvon-Durocher et al. 2014). CH₄ emissions also have a higher temperature dependence than ER, such that the ratio of CH₄ to CO₂ emissions was found to increase markedly with seasonal increases in temperature (Yvon-Durocher et al. 2014). Similarly, we observed a significant increase in the ratio of annual CH₄ to ER along geographic temperature gradients, ranging from 0.4% to 7.9%, with a median value of 2.8% across the dataset (Fig. 6d). This relationship suggests that warming may result in a greater relative contribution of CH₄ to total carbon emissions from wetland ecosystems. With a growing FLUXNET CH₄ database, it will be possible to further explore the dominant controls on CH₄ fluxes within and across ecosystem types, as well as further investigate the temperature dependence of ecosystem-scale CH₄ exchange (Schipper et al. 2014; Arcus et al. 2016; Yvon-Durocher et al. 2014).

Time scales of variability. Methane fluxes exhibited strong variability over a range of time scales, with the

variation across time scales differing between wetland types (Fig. 7). As observed previously (Sturtevant et al. 2016), the seasonal time scale dominated CH₄ flux variability across wetland types, but was most pronounced in rice paddies, which have a distinct growing season, and least pronounced in bogs (Fig. 7). Across ecosystem types, variation was lowest at the multiday scale, although multiday CH₄ flux variation was slightly greater in rice paddies and wet tundra, potentially indicating greater water table fluctuations (Sturtevant et al. 2016), particularly at rice paddy sites, which are subject to seasonal drainage (Knox et al. 2016; Runkle et al. 2019). Whereas some studies report a strong diel pattern in CH₄ emissions from wetlands and rice paddies (Knox et al. 2016; Chu et al. 2014; Morin et al. 2014b; Kim et al. 1999), others have found little or no diel variation (Rinne et al. 2018; Jackowicz-Korczyński et al. 2010; Yagi and Minami 1990; Nadeau et al. 2013). Across wetland types, diel variation was greatest in freshwater marshes (Fig. 7), which is consistent with the observations that the vegetation at sites with a strong diel cycle of CH₄ emissions is typically dominated by species with convective gas flow such as *Phragmites australis* or *Typha* spp. (Brix et al. 1992; Chanton et al. 1993). Bogs, fens, and wet tundra showed the greatest variation at the hourly scale (Fig. 7). This is likely in part due to typically lower fluxes at these sites as hourly perturbations of turbulent time series are largely dominated by noise (Hollinger and Richardson 2005), as well as the fact that near-surface turbulence and short-term pressure fluctuations can strongly influence CH₄ exchange in these peat dominated ecosystems (Nadeau et al. 2013; Sachs et al. 2008).

Gap-filling performance and uncertainty quantification.

The performance of the neural networks varied strongly across sites (Fig. 8). Model r^2 , calculated from the median ANN prediction and observed fluxes at each site, ranged from ~0 to 0.92 across sites, with a median value of 0.41. Across sites, ANN performance was strongly linked to the percentage of total variance at diel and seasonal scales ($r^2 = 0.69$, $p < 0.001$), indicating that across the wide range of observed flux magnitudes, sites with a more distinct seasonal and diel pattern tended to be more predictable (Fig. 8). There was also a significant negative relationship between model r^2 and the percentage of total variance at the hourly scale across sites ($r^2 = 0.72$, $p < 0.001$), because, as noted previously, hourly perturbations are largely dominated by noise (Hollinger and Richardson 2005).

Knowledge of the random errors in half-hourly flux measurements is not only important for

evaluating the uncertainty in cumulative fluxes (e.g., daily, monthly, or annual) and comparing fluxes across tower sites, but it also needed to incorporate information about random flux errors in model-data synthesis activities (Richardson et al. 2006). As noted above, random flux error more closely followed a Laplace rather than Gaussian distribution. Within sites, $\sigma(\delta)$ was not constant, but rather nearly always scaled with the magnitude of CH_4 fluxes (Fig. 9a), as predicted from theory (Richardson et al. 2006). As observed for other fluxes (Richardson et al. 2006), both the slope and intercept of this relationship varied among sites, and depending on the sign of the flux (Fig. 9a). Across sites, random flux error therefore scaled linearly with the magnitude of mean CH_4 flux ($r^2 = 0.86$, $p < 0.001$), even when excluding the two highest CH_4 -emitting sites ($r^2 = 0.46$, $p < 0.001$) (Fig. 9b). Whereas closed-path CH_4 analyzers have been found to have lower random errors and instrument noise compared with open-path sensors (Peltola et al. 2014), there was no clear evidence of a systematic effect of the influence of closed- versus open-path sensors on random errors across sites (Fig. 9).

The total annual cumulative uncertainty in CH_4 fluxes, including both random and gap-filling errors, ranged from ± 0.01 to $\pm 13.4 \text{ g C m}^{-2} \text{ yr}^{-1}$, with a median value of $\pm 1.0 \text{ g C m}^{-2} \text{ yr}^{-1}$ at 95% confidence (Fig. 10a). Relative error decreased exponentially with flux magnitude, ranging from 1.5% to 60% in most cases (Fig. 10b), although a few sites where annual CH_4 sums were near zero had relative errors exceeding 200% (data not shown). The highest relative errors therefore tended to be associated with low CH_4 -emitting sites, such as upland sites and bogs, and the lowest relative errors were generally associated with high CH_4 -emitting sites such as freshwater marshes (Fig. 10b).

FUTURE RESEARCH DIRECTIONS AND NEEDS. Better quantification of CH_4 sources and sinks will improve estimates of regional and global CH_4 budgets and reduce uncertainties in the CH_4 cycle. In this general

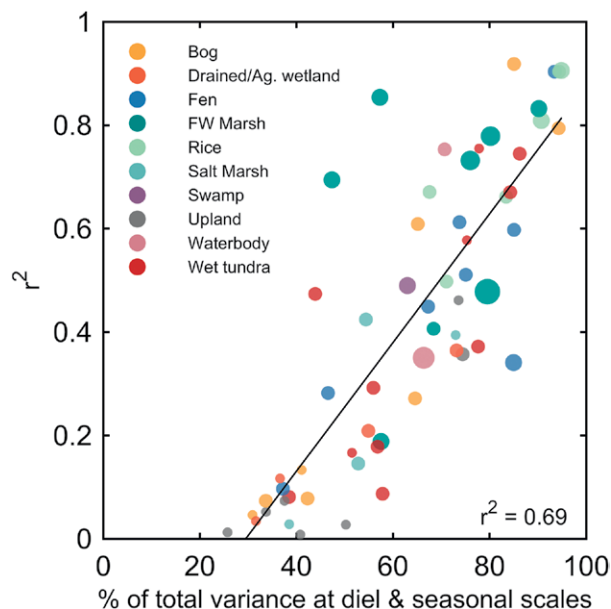


FIG. 8. Relationship between the correlation coefficient (r^2) calculated from the median ANN prediction and observed CH_4 fluxes at each site and the percentage of total variance at diel and seasonal scales ($r^2 = 0.69$, $p < 0.001$). Each site is color coded by ecosystem type. Sizes of the dots are proportional to the magnitude of mean CH_4 flux, where flux magnitude was aggregated into 10 bins for plotting.

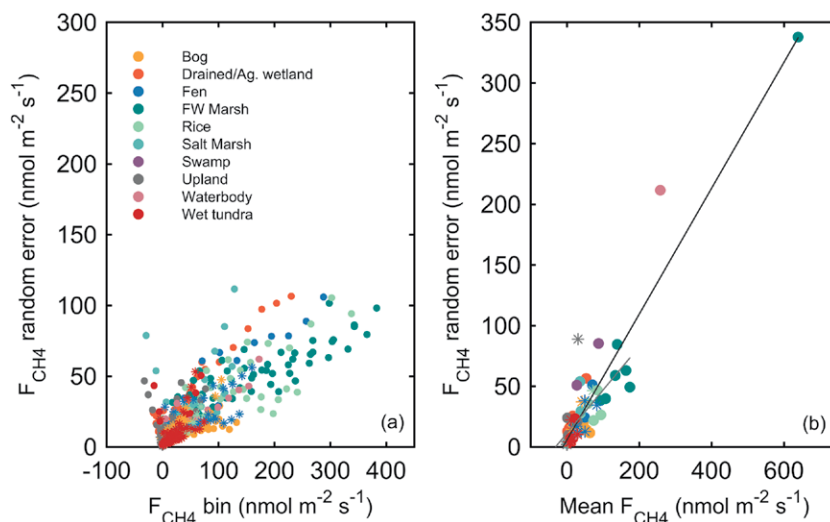


FIG. 9. (a) Scaling of F_{CH_4} random flux measurement error [$\sigma(\delta)$] with flux magnitude for all sites with a significant linear relationship between random error and flux magnitude (95% of all sites). Data at each site were placed into 10 bins (Oikawa et al. 2017). (b) Scaling of F_{CH_4} random flux measurement error, characterized by the standard deviation of the double-exponential distribution [$\sigma(\delta)$], with mean flux magnitude across sites. There was a significant linear relationship between $\sigma(\delta)$ and the magnitude of mean CH_4 flux [$\sigma(\delta) = 0.5 \times F_{\text{CH}_4} + 5.9$, $r^2 = 0.86$, $p < 0.001$], even when excluding the two highest CH_4 -emitting sites [$\sigma(\delta) = 0.4 \times F_{\text{CH}_4} + 11.3$, $r^2 = 0.46$, $p < 0.001$]. Note that circles represent sites with open-path CH_4 analyzers while asterisks represent sites with closed-path sensors.

context, high-frequency observations of ecosystem-scale CH_4 emissions should help constrain bottom-up CH_4 budgets, improve our understanding of the environmental factors controlling CH_4 fluxes, and inform and validate land surface models used to estimate global CH_4 fluxes. Unlike well-established efforts synthesizing CO_2 , water vapor, and energy observations, no such global data synthesis or initiative previously existed for CH_4 . The database presented here addresses this gap with the EC community by organizing the collection and aggregation of a global EC CH_4 database through FLUXNET.

EC flux data quality assessment.

Much of what has been learned within FLUXNET for CO_2 , water vapor, and energy measurements is informing, and should continue to inform, new efforts for CH_4 . Reliable EC measurements of CO_2 and water vapor fluxes have been conducted at hundreds of sites across broad regional networks (Papale et al. 2012), and substantial efforts have focused on developing best practices and harmonizing approaches across sites to ensure consistent, high-quality flux measurements (Aubinet et al. 1999; Reichstein et al. 2005; Moffat et al. 2007). CH_4 fluxes are often characterized by small fluxes with episodic spikes, and additional research is needed to ensure reliable measurements (Peltola et al. 2014, 2013), and refine and standardize methods and routines for data processing and quality checking (Nemitz et al. 2018; Schaller et al. 2018). Recent efforts provided guidance on instrument selection, setup and maintenance, and data processing for EC CH_4 flux measurements (Nemitz et al. 2018). However, with respect to instrument setup and data processing, more research is needed in best practices for storage flux quantification, despiking, and u_c filtering (Nemitz et al. 2018).

Gap-filling. Whereas neural networks have shown strong performance for gap-filling CH_4 fluxes (Dengel et al. 2013; Knox et al. 2016), our results reveal some of the challenges of gap-filling CH_4 fluxes at sites with low fluxes and/or a lack of seasonal and diel variation (Fig. 8). More research is therefore needed

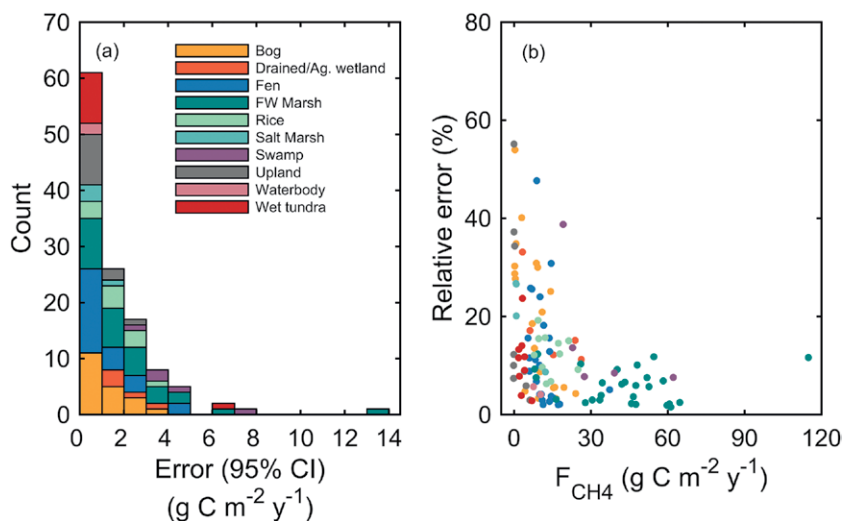


FIG. 10. (a) Histogram of total random error ($\text{g C m}^{-2} \text{yr}^{-1}$) in annual CH_4 flux at 95% confidence, where count refers to the number of site years of measurements. The cumulative gap-filling and random measurement uncertainties of gap-filled and original values were added in quadrature to estimate the total random uncertainty at each site. (b) Relationship between annual CH_4 flux ($\text{g C m}^{-2} \text{yr}^{-1}$) and relative error (i.e., total random error divided by flux magnitude; %).

for best practices for gap-filling to estimate annual CH_4 budgets (Nemitz et al. 2018). For example, there has yet to be a comprehensive analysis comparing a wide range of gap-filling approaches for CH_4 fluxes similar to the study by Moffat et al. (2007) for CO_2 exchange. While ANNs are one gap-filling method (Dengel et al. 2013; Shoemaker et al. 2014; Morin et al. 2014a), numerous other gap-filling approaches exist, including nonlinear regression techniques, mean diurnal variation, lookup tables, marginal distribution sampling, and the multiple imputation method (Moffat et al. 2007; Vitale et al. 2019). Future efforts should focus on systematically investigating these approaches across a range of sites to provide best practices for gap-filling CH_4 exchange.

Ancillary measurements. Along with research that addresses the challenges of measuring and processing EC CH_4 fluxes, key ancillary variables to help gap-fill, predict, and scale CH_4 fluxes should also be measured more comprehensively across sites. For instance, although WTD is known to strongly influence CH_4 emissions (Turetsky et al. 2014; Treat et al. 2018b), as noted above, only half of the sites currently included in the database report water table position. Generally, EC CH_4 measurements are implemented at sites also collecting CO_2 fluxes and common meteorological measurements used in the flux community; however, guidelines are only

beginning to emerge for which additional supporting variables should be collected at sites measuring CH_4 fluxes (Nemitz et al. 2018).

Measurements of variables beyond those relevant for CO_2 are needed to better understand and predict the complex and interacting processes of CH_4 production, consumption, and transport, the latter of which includes diffusion, ebullition, and plant-mediated transport. Guidance on the description of some basic variables affecting these processes is available through new protocols in the flux community detailing soil meteorological measurements, ancillary vegetation measurements, and site description, management and disturbance (Saunders et al. 2018; Op De Beeck et al. 2018; Gielen et al. 2018). These protocols provide guidance on variables such as soil temperature and soil moisture profiles, water table depth and snow depth, soil pH and soil type, bulk density, and livestock density. However, although WTD is an easily measured proxy for anaerobic conditions, direct and continuous measurement of redox potential and oxygen content in particular would be valuable additional measurements (Nemitz et al. 2018). Similarly, measuring variables such as conductivity, below-ground CH_4 concentrations, dissolved organic carbon concentrations, and the presence of alternative electron acceptors such as nitrate, iron, sulfate, and humic substances in the water and soil column would provide useful information for the interpretation of CH_4 emissions. Stable isotope analyses of CH_4 are also valuable as they provide important information on mechanisms of CH_4 production, transport, and oxidation (Chanton et al. 1997; Marushchak et al. 2016). Detailed information on soil microbial communities driving CH_4 production and consumption could also be helpful (Kwon et al. 2017). Vegetation biomass, species composition, and phenology are also important variables to consider, because plants are a primary source of carbon substrates for methanogenic metabolism, and they mediate CH_4 transport through aerenchymous tissue (Kwon et al. 2017; Joabsson et al. 1999; Carmichael et al. 2014). New guidance is now available for such measurements at flux tower locations (Gielen et al. 2018; Hufkens et al. 2018). Continuing to develop a consensus on best practices for ancillary measurements is important for interpreting, gap-filling, and upscaling CH_4 flux measurements.

Characterizing spatial variability. CH_4 fluxes exhibit finescale spatial variability that can span orders of magnitude within a landscape (Peltola et al. 2015; Marushchak et al. 2016; Desai et al. 2015; Treat et al.

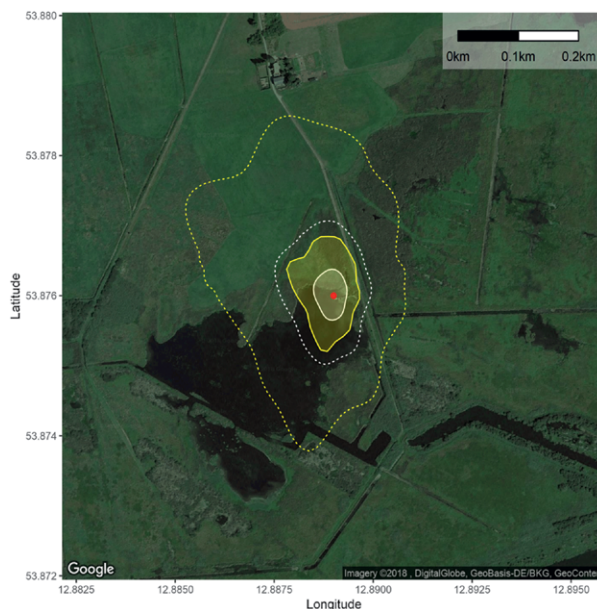


FIG. 11. Footprint climatology for a eutrophic shallow lake on a formerly drained fen in Germany (Zarnekow; DE-Zrk) illustrating the importance of footprint analysis for the interpretation of EC measurements of CH_4 . Here we used two footprint models, including the model of Kormann and Meixner (2001) (yellow) and Kljun et al. (2015) (white). The footprint climatology was calculated by aggregating all half-hour footprints within a year. The dashed lines enclose the areas aggregating to 80% of source areas, while solid lines enclose the 50% of source areas.

2018a; Iwata et al. 2018), attributable to heterogeneous soil properties and moisture conditions, vegetation composition, and land use (Davidson et al. 2016; Parmentier et al. 2011; Chamberlain et al. 2018). Furthermore, there is evidence that traditionally unmeasured surfaces (i.e., tree stems) are important sources of CH_4 to the atmosphere and could explain spatial heterogeneity within ecosystems (Barba et al. 2019). Accurately representing spatial heterogeneity and the relative fraction of uplands and wetlands is imperative for interpreting and predicting CH_4 emissions within many ecosystems, and for upscaling flux measurements regionally and globally as wetlands are hot spots for carbon cycling (Treat et al. 2018a; Tuovinen et al. 2019; Rößger et al. 2019). Flux footprint analysis characterizing the fractional coverage of the dominant surface types, particularly the fraction of open water and aerenchymatous plants, is important for interpreting EC CH_4 flux measurements and quantifying annual CH_4 budgets at spatially heterogeneous sites (Franz et al. 2016; Helbig et al. 2017a; Jammert et al. 2017) (Fig. 11). This integration can be achieved by combining CH_4 measurements, flux footprint analysis, and near-surface

TABLE A1. The ecosystem type is based on the classification of Olefeldt et al. (2013) and Treat et al. (2018b). Biome was based on the classification of Olson et al. (2001) and extracted using site coordinates. Vegetation type was based on the International Geosphere-Biosphere Programme (IGBP) definition. Salinity regime includes freshwater (FW) or saltwater (SW) wetlands. Disturbance is based on the classification of Turetsky et al. (2014). Data from all sites are publicly available, primarily through AmeriFlux and the European Database Cluster, and in a few cases, through other databases/repositories. Site DOIs are specified where applicable.

Site ID	Site name	Country	Lat (°N)	Lon (°E)	Biome	IGBP
US-ICs	Wet sedge tundra	U.S.	68.606	-149.311	Tundra	WET
SE-St1	Stordalen palsland (mire)	Sweden	68.350	19.050	Tundra	WET
SE-Sto	Stordalen Palsa bog	Sweden	68.356	19.050	Tundra	WET
RU-Vrk	Seida/Vorkuta	Russia	67.055	62.940	Tundra	WET
RU-Ch2	Chersky reference	Russia	68.617	161.351	Tundra	WET
RU-Che	Chersky	Russia	68.613	161.341	Tundra	WET
RU-SAM	Samoylov	Russia	72.374	126.496	Tundra	WET
US-NGB	NGEE Barrow	U.S.	71.280	-156.609	Tundra	WET
US-Beo	Barrow	U.S.	71.281	-156.612	Tundra	WET
US-Bes	Barrow	U.S.	71.281	-156.596	Tundra	WET
US-Atq	Atkasuk	U.S.	70.470	-157.409	Tundra	WET
US-lvo	Ivotuk	U.S.	68.486	-155.750	Tundra	WET
—	Black spruce forest	U.S.	64.700	-148.320	Boreal forests/taiga	ENF
—	Rich fen	U.S.	64.704	-148.313	Boreal forests/taiga	WET
—	Thermokarst collapse scar bog	U.S.	64.700	-148.320	Boreal forests/taiga	WET
FI-Lom	Lompolojankka	Finland	67.997	24.209	Boreal forests/taiga	WET
SE-Deg	Degero	Sweden	64.182	19.557	Boreal forests/taiga	WET
CA-SCC	Scotty Creek—Peat plateau/collapse scar	Canada	61.308	-121.299	Boreal forests/taiga	ENF
CA-SCB	Scotty Creek bog	Canada	61.309	-121.299	Boreal forests/taiga	WET
US-NGC	NGEE Arctic Council	U.S.	64.861	-163.701	Boreal forests/taiga	WET
US-Uaf	University of Alaska Fairbanks	U.S.	64.866	-147.856	Boreal forests/taiga	WET
FI-Sii	Siikaneva I (FI-Sii)	Finland	61.833	24.193	Boreal forests/taiga	WET
FI-Si2	Siikaneva II	Finland	61.837	24.170	Boreal forests/taiga	WET
US-Myb	Mayberry wetland	U.S.	38.050	-121.765	Temperate	WET
US-Sne	Sherman Island restored wetland	U.S.	38.037	-121.755	Temperate	WET
US-Tw1	Twitchell west pond wetland	U.S.	38.107	-121.647	Temperate	WET
US-Tw4	Twitchell east end wetland	U.S.	38.103	-121.641	Temperate	WET
US-Twt	Twitchell rice	U.S.	38.109	-121.653	Temperate	CRO - Rice
US-Bi2	Bouldin Island corn	U.S.	38.109	-121.535	Temperate	CRO - Other
US-Bi1	Bouldin Island alfalfa	U.S.	38.102	-121.504	Temperate	CRO - Other
US-Snd	Sherman Island	U.S.	38.037	-121.754	Temperate	CRO - Other
US-OWC	Old Woman Creek	U.S.	41.380	-82.512	Temperate	WET
US-Orv	Olentangy River Wetland Research Park	U.S.	40.020	-83.018	Temperate	WET
NZ-Kop	Koputai	New Zealand	-37.388	175.554	Temperate	WET
IT-Cas	Castellaro	Italy	45.070	8.718	Temperate	CRO - Rice
US-WPT	Winous Point north marsh	U.S.	41.465	-82.996	Temperate	WET
US-CRT	Curtice Walter-Berger cropland	U.S.	41.628	-83.347	Temperate	CRO - Other
US-Los	Lost Creek	U.S.	46.083	-89.979	Temperate	WET
JP-Mse	Mase paddy flux site (MSE)	Japan	36.054	140.027	Temperate	CRO - Rice
JP-Swl	Suwa Lake site	Japan	36.047	138.108	Temperate	WAT
IT-BCi	Borgo Cioffi	Italy	40.524	14.957	Temperate	CRO - Other
—	Hongyuan	China	32.800	102.550	Temperate	GRA
US-NC4	NC Alligator River	U.S.	35.788	-75.904	Temperate	WET
DE-SfN	Schechenfilz Nord	Germany	47.806	11.328	Temperate	WET
US-Ho1	Howland Forest (main tower)	U.S.	45.204	-68.740	Temperate	ENF
US-HRA	Humnoke farm rice field AWD, United States	U.S.	34.585	-91.752	Temperate	CRO - Rice
US-HRC	Humnoke farm rice field conventional, United States	U.S.	34.589	-91.752	Temperate	CRO - Rice
KR-CRK	Cheorwon rice paddy	South Korea	38.201	127.251	Temperate	CRO - Rice
DE-Zrk	Zarnkow	Germany	53.876	12.889	Temperate	WET
DE-Dgw	Dagowsee	Germany	53.151	13.054	Temperate	WAT
US-MRM	Marsh Resource Meadowlands Mitigation Bank	U.S.	40.816	-74.044	Temperate	WET
—	Bog Lake peatland	U.S.	47.530	-93.470	Temperate	WET
—	MacArthur Agro-Ecology Research Center	U.S.	27.163	-81.187	Temperate	CRO - Other
JP-BBY	Bibai bog	Japan	43.323	141.811	Temperate	WET
US-Stj	St. Jones Reserve	U.S.	39.088	-75.437	Temperate	WET
US-Srr	Suisun marsh—Rush Ranch	U.S.	38.201	-122.026	Temperate	WET
AT-Neu	Neustift	Austria	47.117	11.318	Temperate	GRA
US-LA2	Salvador WMA freshwater marsh	U.S.	29.859	-90.287	Tropical and subtropical	WET
US-LA1	Pointe-aux-Chenes brackish marsh	U.S.	29.501	-90.445	Tropical and subtropical	WET
MY-MLM	Maludam	Malaysia	1.454	111.149	Tropical and subtropical	WET

Wetland				
Ecosystem type	Salinity	disturbance	Site PI	Data DOI/location
Wet tundra	FW	Undisturbed	Eugenie Euskirchen	DOI:10.17190/AMF/1246130
Fen	FW	Undisturbed	Thomas Friborg	European Fluxes Database Cluster
Bog	FW	Undisturbed	Thomas Friborg	European Fluxes Database Cluster
Wet tundra	FW	Undisturbed	Thomas Friborg	European Fluxes Database Cluster
Wet tundra	FW	Undisturbed	Mathias Goeckede	European Fluxes Database Cluster
Wet tundra	FW	Drying	Mathias Goeckede	European Fluxes Database Cluster
Wet tundra	FW	Undisturbed	Torsten Sachs	European Fluxes Database Cluster
Wet tundra	FW	Undisturbed	Margaret Torn	DOI:10.17190/AMF/1436326
Wet tundra	FW	Undisturbed	Donatella Zona	AmeriFlux
Wet tundra	FW	Undisturbed	Donatella Zona	AmeriFlux
Wet tundra	FW	Undisturbed	Donatella Zona	DOI:10.17190/AMF/1246029
Wet tundra	FW	Undisturbed	Donatella Zona	DOI:10.17190/AMF/1246067
Upland	—	—	Eugenie Euskirchen	www.lter.uaf.edu/data/data-detail/id/708
Fen	FW	Undisturbed	Eugenie Euskirchen	www.lter.uaf.edu/data/data-detail/id/708
Bog	FW	Undisturbed	Eugenie Euskirchen	www.lter.uaf.edu/data/data-detail/id/708
Fen	FW	Undisturbed	Annalea Lohila	European Fluxes Database Cluster
Fen	FW	Undisturbed	Matthias Peichl, Mats Nilsson	European Fluxes Database Cluster
Peat plateau	FW	—	Oliver Sonnentag	DOI:10.17190/AMF/1480303
Bog	FW	Undisturbed	Oliver Sonnentag	AmeriFlux
Wet tundra	FW	Undisturbed	Margaret Torn	AmeriFlux
Bog	FW	Undisturbed	Masahito Ueyama	DOI:10.17190/AMF/1480322
Fen	FW	Undisturbed	Timo Vesala, Ivan Mammarella	European Fluxes Database Cluster
Bog	FW	Undisturbed	Timo Vesala, Ivan Mammarella	European Fluxes Database Cluster
Marsh	FW	Wetting	Dennis Baldocchi	DOI:10.17190/AMF/1246139
Marsh	FW	Wetting	Dennis Baldocchi	DOI:10.17190/AMF/1418684
Marsh	FW	Wetting	Dennis Baldocchi	DOI:10.17190/AMF/1246147
Marsh	FW	Wetting	Dennis Baldocchi	DOI:10.17190/AMF/1246148
Rice	FW	—	Dennis Baldocchi	DOI:10.17190/AMF/1246151
Drained/agricultural wetland	FW	Drying	Dennis Baldocchi	DOI:10.17190/AMF/1419513
Drained/agricultural wetland	FW	Drying	Dennis Baldocchi	DOI:10.17190/AMF/1480317
Drained/agricultural wetland	FW	Drying	Dennis Baldocchi	DOI:10.17190/AMF/1246094
Marsh	FW	Undisturbed	Gil Bohrer	DOI:10.17190/AMF/1246094
Marsh	FW	Undisturbed	Gil Bohrer	DOI:10.17190/AMF/1246135
Bog	FW	Undisturbed	Dave Campbell	https://researchcommons.waikato.ac.nz/handle/10289/11393
Rice	FW	—	Alessandro Cescatti	European Fluxes Database Cluster
Marsh	FW	Wetting	Jiquan Chen, Housen Chu	DOI:10.17190/AMF/1246155
Upland	—	—	Jiquan Chen, Housen Chu	DOI:10.17190/AMF/1246156
Fen	FW	Undisturbed	Ankur Desai	DOI:10.17190/AMF/1246071
Rice	FW	—	Akira Miyata	European Fluxes Database Cluster
Waterbody	FW	Undisturbed	Hiroki Iwata	European Fluxes Database Cluster
Upland	—	—	Vincenzo Magliulo	European Fluxes Database Cluster
Upland	—	—	Shuli Niu	European Fluxes Database Cluster
Swamp	FW	Undisturbed	Asko Noormets	DOI:10.17190/AMF/1480314
Bog	FW	Undisturbed	Hans Peter Schmid	European Fluxes Database Cluster
Upland	—	—	Andrew Richardson	DOI:10.17190/AMF/1246061
Rice	FW	—	Benjamin Runkle	AmeriFlux
Rice	FW	—	Benjamin Runkle	AmeriFlux
Rice	FW	—	Youngryel Ryu, Minseok Kang	European Fluxes Database Cluster
Fen	FW	Wetting	Torsten Sachs	European Fluxes Database Cluster
Waterbody	FW	Undisturbed	Torsten Sachs	European Fluxes Database Cluster
Salt marsh	SW	Wetting	Karina Schäfer	AmeriFlux
Fen	FW	Undisturbed	Shahi Verma	AmeriFlux
Drained/agricultural wetland	FW	Drying	Jed Sparks, Samuel Chamberlain	AmeriFlux
Bog	FW	Undisturbed	Masahito Ueyama	European Fluxes Database Cluster
Salt marsh	SW	Undisturbed	Rodrigo Vargas	DOI:10.17190/AMF/1480316
Salt marsh	SW	Undisturbed	Lisamarie Windham-Myers	DOI:10.17190/AMF/1418685
Upland	—	—	Georg Wohlfahrt	European Fluxes Database Cluster
Marsh	FW	Undisturbed	Ken Krauss	AmeriFlux
Salt marsh	SW	Undisturbed	Ken Krauss	AmeriFlux
Swamp	FW	Undisturbed	Angela Tang	DOI:10.5281/zenodo.1161966

(e.g., phenocams) and/or high-resolution drone or satellite remote sensing data, and should be common practice for all sites measuring CH₄ fluxes.

Spatial variability in ecosystem-scale CH₄ flux can further be examined by combining chamber and EC measurements, including manual and autochambers, multitower approaches, and airborne flux measurements (Peltola et al. 2015; Zona et al. 2016; Helbig et al. 2017a; Wolfe et al. 2018; Kohnert et al. 2018; Lai et al. 2014; McNicol et al. 2017). Integrating additional observations such as information on microbial communities, isotopic measurements, and laboratory incubation observations along with chamber and EC CH₄ flux measurements can further help explain CH₄ dynamics across scales (Angle et al. 2017; Chamberlain et al. 2018; Yang et al. 2017). However, as discussed above, additional research is needed to reconcile differences in fluxes measured across scales (Gioli et al. 2004; Holm et al. 2016; Meijide et al. 2011). Explicitly considering source area composition and spatial heterogeneity will provide enhanced processed-based understanding of CH₄ fluxes and improve upscaled regional and global estimates of CH₄ emissions, which can help reconcile the discrepancy between bottom-up and top-down budgets (Saunois et al. 2016a; Morin et al. 2017; Davidson et al. 2016).

More sites in key regions. We expect the number of flux towers measuring CH₄ fluxes will continue to grow (Chu et al. 2017; Pastorello et al. 2017; Morin 2018), but our compilation of EC CH₄ flux sites highlights key underrepresented regions where future flux towers are needed or where more efforts are needed for existing but nonreporting towers to contribute to FLUXNET (Fig. 1). As noted previously, notable gaps include both tropical and subtropical regions, as well as eastern Canada, and the boreal forests of Russia. Figure 1 also provides guidance on where new towers could be strategically located to help reconcile differences between top-down and bottom-up estimates of wetland CH₄ emissions. In particular, substantial disagreements between top-down and bottom-up estimates are found over the Congo basin, the Inner Niger delta, the Orinoco River delta, the Marañón–Ucayali palm swamps, the Pantanal, the Ganges–Brahmaputra delta, Sumatra, the western Siberian lowlands, and the Hudson Bay lowlands (Fig. 1). However, the placement of new towers is a strong function of the scientific question being asked and research funding priorities, and therefore the optimal tower network could be different for different applications (Mahecha et al. 2017; Papale et al. 2015; Villarreal et al. 2018).

Better understanding and representing processes. One of the biggest challenges for understanding ecosystem functioning is resolving overlapping, asynchronous (i.e., lagged) and nonlinear processes (Sturtevant et al. 2016). This challenge is particularly relevant for interpreting continuous, ecosystem-scale measurements of CH₄ exchange where scale-specific, nonlinear, and lagged processes may dominate (Franz et al. 2016; Sturtevant et al. 2016; Knox et al. 2018). For instance, CH₄ emission responses to water table fluctuation can be nonlinear and lagged on the order of days to months (Goodrich et al. 2015; Sturtevant et al. 2016). CH₄ flux has also been observed to lag GPP by hours to days (Rinne et al. 2018; Hatala et al. 2012). Adequately representing these dynamics in process models is important, and further research is needed to better characterize the complex and nonlinear processes influencing ecosystem-scale CH₄ exchange across time scales.

The complex nature of CH₄ flux dynamics requires moving beyond traditional linear correlation and regression, and using methods such as wavelets, information theory, and Granger causality that are more tailored to address scale, nonlinearity, and lags directly (Stoy et al. 2005; Vargas et al. 2011; Schäfer et al. 2014; Knox et al. 2016; Detto et al. 2012). Through a USGS Powell Center working group activity, we will continue to investigate controls on CH₄ emissions within and across wetland types. To further explore interactions between ecosystem-scale CH₄ exchange and drivers across time scales, wavelet analysis will be combined with information theory to explore biosphere–atmosphere interactions regardless of form or asynchrony (Sturtevant et al. 2016; Knox et al. 2018; Chamberlain et al. 2018). By coupling wavelet decomposition with information theory, future research will investigate key controls on CH₄ fluxes across time scales, as well as the importance of nonlinearities and lags in predicting CH₄ flux dynamics. Future research will also use the global CH₄ database to parameterize and benchmark the performance of land surface models of global CH₄ emissions, providing a unique opportunity for informing and validating biogeochemical models.

Coordinating, organizing and improving the integration of CH₄ fluxes in regional networks and ultimately FLUXNET will bring us one step closer to achieving the goal of providing flux information “everywhere and all of the time” (Baldocchi 2008). In the long term, we hope to integrate the global eddy covariance CH₄ database with other methods for measuring CH₄ fluxes, such as chamber, aircraft, and satellite measurements. By integrating CH₄ flux

measurements, remote sensing, and modeling, we aim to better characterize CH₄ emissions from terrestrial ecosystems and ultimately reduce uncertainties in the global CH₄ cycle.

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APPENDIX: SITES CURRENTLY INCLUDED IN THE DATABASE. Table A1 presents characteristics of sites currently included in the database.

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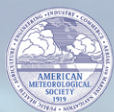
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