

An expert survey on chamber measurement techniques and data handling procedures for methane fluxes

Katharina Jentsch^{1,2}, Lona van Delden¹, Matthias Fuchs³, Claire C. Treat^{1,4}

¹Alfred Wegener Institute (~~AWI~~) Helmholtz Centre for Polar and Marine Research, Potsdam, Germany

²Institute of Environmental Science and Geography, University of Potsdam, Potsdam, Germany

³Renewable and Sustainable Energy Institute, University of Colorado Boulder, Boulder, CO, USA

⁴Department of Agroecology, Aarhus University, Denmark,

Correspondence to: Katharina Jentsch (katharina.jentsch@awi.de)

Abstract.

Methane is an important greenhouse gas but the magnitude of global emissions in particular from natural sources remains highly uncertain. To estimate methane emissions on large spatial scales, methane flux data sets from field measurements collected and processed by many different researchers must be combined. One common method for obtaining in-situ methane flux measurements are flux chambers. We hypothesize that considerable uncertainty might be introduced into ~~such~~ data synthesis products derived from chamber measurements due to by the many different the variety of measurement setups, data processing and quality control approaches used ~~to collect, process and quality control chamber measurements of methane fluxes~~ within the chamber flux community. Existing guidelines on chamber measurements promote more standardized measurement and data processing techniques but to our knowledge, so far, no study has investigated which methods are actually used within the chamber flux community. Therefore, we aimed to identify the key discrepancies~~major differences~~ between the measurement and data handling procedures ~~used~~implemented ~~approaches~~ for chamber methane fluxes ~~used~~ by different researchers.

We conducted an expert survey to collect information on why, where, and how scientists conduct chamber-based methane flux measurements and how they handle the resulting data, including field sites, research questions, measurement setups and routines as well as data processing and quality control of data. We received 36 responses from researchers in North America, Europe, and Asia which ~~revealed~~indicated that 80% of respondents have adopted high-frequency, multi-gas analyzers to obtain high-frequency (< 1 Hz) methane concentration measurements over a total chamber closure time of typically with most measurement times falling between 2 and 5 minutes. Most but not all of the respondents use recommended chamber designs, including features such as airtight sealing, fans, and a pressure vent. We ~~asked about the participants' approach to quality control and~~ presented a standardized set of methane concentrations timeseries recorded during from observed flux chamber measurements and derived CH₄ flux estimates based on the processing and quality control approaches suggested by the survey participants, then included this information for flux calculations. The responses showed broad disagreement among the experts ~~on concerning the~~ processes that they consider responsible for ~~resulting in~~ nonlinear methane concentration increases.

Formatted: English (United States)

Formatted: Line spacing: Multiple 1,15 li

Formatted: English (United States), Superscript

Formatted: English (United States)

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: Subscript

Furthermore, there was a tendency to discard low or negative CH₄ fluxes. Based on the expert responses, we estimated a variability-uncertainty of 28% introduced by different researchers deciding differently on discarding vs. accepting a measurement when processing a representative data set of chamber measurement. Different researchers choosing different time periods within the same measurement for flux calculation caused an additional variability-uncertainty of 17%. Our study highlights the importance need to understand which the processes are causing drivers of the patterns in CH₄ concentrations visible from high-resolution analyzers and well as the need for standardized data handling procedures in and guidelines for future chamber methane flux measurements. This is highly important to reliably quantify methane fluxes all over the world.

The survey results as well as the questionnaire are publicly available at <https://doi.org/10.1594/PANGAEA.971695> (Jentsch et al., 2024b).

1 Introduction

Methane (CH₄) is an important greenhouse gas with 45 times the global warming potential of carbon dioxide (CO₂) on a 100-year timescale (Neubauer, 2021). However, emission estimates differ largely between “top-down” atmospheric measurement inversions and “bottom-up” approaches using data-constrained or process-based models (Kirschke et al., 2013; Saunois et al., 2020). Natural emissions, especially bottom-up estimates of wetland emissions, are the largest source of uncertainty to the global CH₄ budget due to the poorly constrained areal extent of wetlands and other methane-producing ecosystems like lakes, streams, and reservoirs, highly uncertain CH₄ process parameterization, and a lack of validation data sets (Melton et al., 2013; Saunois et al., 2020).

Despite more than thirty years of chamber-based methane flux measurements from wetland ecosystems (Bartlett & Harriss, 1993; Harriss et al., 1985), developing large-scale methane validation data sets remain challenging. One approach to obtain large-scale validation data sets for CH₄ fluxes has been to create synthesis datasets of measurements collected made by multiple researchers using chamber-based methane flux measurements (Kuhn et al., 2021; Olefeldt et al., 2013; Treat et al., 2018). The advantage of using the closed-chamber technique over in-situ measurements operating on larger spatial scales is that the resulting data sets data sets should can capture the high spatial and temporal variability in natural CH₄ emissions with small-scale spatial changes in environmental and ecological conditions (Frenzel and Karofeld, 2000; Laine et al., 2007; Moore and Knowles, 1990; Waddington and Roulet, 1996). This is commonly achieved with the When applying the closed-chamber technique, a in which a chamber is placed on top of the soil and the change in gas concentrations in the chamber headspace is monitored over time to estimate the exchange of CH₄ between soil, plants, and atmosphere on the microscale (e.g. Livingston and Hutchinson, 1995). The rate of change in gas concentrations, after correcting for temperature and pressure conditions using the ideal gas law, is then used to compute the flux of CH₄ through the surface area covered by the chamber (Holland et al.,

Formatted: Subscript

Formatted: Subscript

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: Font: (Default) Times New Roman

Formatted: Indent: First line: 0 cm

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

1999). However, despite more than thirty years of chamber-based methane flux measurements from wetland ecosystems (Bartlett & Harriss, 1993; Harriss et al., 1985), developing large-scale methane validation data sets remains challenging.

Two approaches are typically used for measuring the CH₄ methane gas concentrations inside the chamber are typically used: manual sampling and in-line gas analyzers. Manual sampling of gas concentrations involves extracting gas samples from the chamber headspace in regular time intervals using syringes and subsequently analyzing them for CH₄ concentrations on a gas chromatograph. A linear fit is then usually applied to the CH₄ concentration measurements over time and its slope is used as the flux estimate after correction for the pressure and temperature inside the chamber (Holland et al., 1999). Manual sampling of the chamber headspace is typically characterized by a low sampling frequency which requires a relatively long chamber closure time. The considerations here are balancing the time needed to get a detectable change in CH₄ concentrations versus shorter measurement times to reduce chamber effects (Holland et al., 1999).

With the advances in laser spectroscopy, manual sampling is increasingly replaced by continuously circling chamber air through an in-line gas analyzer which performs high-frequency (>1 Hz), high-accuracy, real-time measurements of the CH₄ concentration. Through their portability and with reduced measurement times, such multigas analyzers have opened new possibilities, particularly for the analysis of key trace gases like CH₄ and N₂O. At the same time, the high frequency and high accuracy of the concentration measurements uncover chamber-induced artefacts and events of ebullitive CH₄ emission that are superimposed on the signal of natural diffusion of CH₄ between soil, plants, and atmosphere. Leakage of gas from the chamber (Hutchinson and Livingston, 2001), a saturation effect changing the concentration gradient between soil and chamber headspace over time (Livingston and Hutchinson, 1995), and natural CH₄ ebullition (Strack et al., 2005) as well as ebullition triggered by the chamber placement can all lead to a deviation of the concentration change from the linear increase expected for a constant diffusive flux. These observations call for a reassessment of the measurement, processing, and quality control (QC) approaches to minimize the influence of chamber effects on the flux estimates.

Besides the general lack of validation data sets, existing data sets that combine flux data collected by different researchers likely include additional uncertainty due to the variety of measurement and data handling/processing approaches used. Several studies have assessed the difference in flux estimates resulting from different chamber setups (Pihlatie et al., 2013; Pumpanen et al., 2004) and from different data processing approaches such as using nonlinear as compared to linear fits to the gas concentration measurements over time (Forbrich et al., 2010; Healy et al., 1996; Pirk et al., 2016). Such experimental and modelling studies have contributed to several guidelines for chamber measurements that were published in an attempt to establish a more standardized protocol for flux measurements. These best-practice guidelines for chamber measurements summarize recommendations on chamber designs (e.g., Clough et al., 2020) as well as on the entire workflow from measurement to data processing and quality control (e.g., de Klein and Harvey, 2012; Fiedler et al., 2022; Maier et al., 2022).

While guidelines outlining best measurement practices for chamber measurements provide a well-founded summary of methods recommended to collect high-quality flux data, they cannot control the approaches that are actually used by the individual researchers. Discussions with other researchers who use chambers to measure methane fluxes revealed that a wide range of measurement and processing approaches are still being used by different researchers. At the same time, method

Formatted: English (United States)

Formatted: Font: 10 pt, English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

descriptions for chamber-based flux data sets collection are often lacking detailed metadata reports on reporting on the chamber design, flux calculation and QC methods.⁵ This which introduces substantial uncertainty to makes a comprehensive comparisons of chamber-based data sets highly uncertain.

Given that the measures recommendations outlined in guidelines for chamber measurements have significant effects on the magnitude of CH₄ fluxes measured, we need to know how widely implemented these recommendations are and where key differences and knowledge gaps remain. Gathering scientific and technical informationsights from experts is necessary to move beyond established theoretical knowledge and can offer further evidence to aid in decision-making (Morgan, 2014). Several studies have recently used expert assessments to gain valuable insights into topicalcurrent climate-change-related

issuestopics (Macreadie et al., 2019; Rosentreter et al., 2024; Schuur et al., 2013). In this study, we use expert judgement derived from a questionnaire to identify the methods for chamber measurements, processing, and QC of CH₄ fluxes that are actually currently used within the flux community and to assess resulting variability and uncertainties.

This study aims to derive starting points for improving the usability of chamber CH₄ flux data sets for large scale synthesis studies through reducing identify the main the discrepancies between the measurement and data handling approaches used within the chamber flux community as identified from an expert surveyby different researchers and their potential effect on the resulting flux data sets. Our objectives were to (1) provide an overview of the chamber designs, measurement setups and routines, flux calculation and QC approaches that are currently used by scientists within the community to quantifystimate CH₄ fluxes as extracted from an expert survey; (2) estimate the variability that is introduced into CH₄ flux data sets by the variety of data handling approaches the variance in QC approaches between different researchers when processing a representative data set of chamber measurements is processed by different researchers; (3) identify major sources of uncertainty resulting from the variety of measurement, calculation, and QC approaches used within the flux community. Our study raises awareness for differences in chamber methods used within the flux community – a potentially considerable but often neglected source of error in synthesis studies that combine flux data sets collected and processed by different researchers. Through identifying major sources of uncertainty resulting from the variety of measurement, calculation, and QC approaches used within the chamber flux community, we derive starting points for eliminating such error sources and rendering individual flux data sets more comparable and combinable and thus better suited for larger scale synthesis studiesand provides a starting point to reduce uncertainties and differences in chamber methods used within the flux community – a potentially considerable but often neglected source of error in synthesis studies that combine flux data sets collected and processed by different researchers.

2 Methods

For this study, we conducted and evaluated an expert survey conducted in 2023 that consisted of two parts – the first part asking questions about the professional background of the participants and the field sites as well as the measurement, calculation and QC approaches that they use for their own chamber measurements of CH₄ fluxes and the second part being an exercise on visual QC of a given set of chamber measurements.

Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

Experts were required to have a minimum expertise of one field season of chamber measurements of CH₄ fluxes.
130 ~~They~~Experts were solicited using emails and conference poster presentations through professional networks ~~using emails and~~
~~conference poster presentations~~, including the Permafrost Carbon Network-~~(pen.org)~~, C-Peat network, ICOS, and through
identification of experts not represented in these networks to increase the number and geographic background of the
participants. Altogether, 46 experts were contacted viaby email. To capture the variety of chamber applications and methods
used within the community, we selected the survey participants to be rather independent from each other in their choice of
135 measurement and data handling approaches.

The survey was estimated to take 40 minutes to complete and the survey language was English. The survey was
administered using LimeSurvey, (Community Edition Version 5.6.68+240625). Survey participants were asked if they wished
to be acknowledged or remain anonymous. Survey participation was voluntary and was not compensated. The survey has been
legally checked by a data protection officer to comply with the EU data protection regulation and involved a privacy policy
140 statement explaining the use and processing of the collected data that needed to be approved by every survey participant prior
to participation. The complete, archived survey can be found in Supplement 1questionnaire and the survey responses are
provided in Jentzsch et al. (2024b).

2.1 Methods of: Survey part 1 – The survey participants and their chamber measurements

In the first, informative part of the survey, we gathered information on the measurement, data processing and QC
145 approaches that the participants use for their own chamber measurements. For this part of the survey we chose a combination
of 20 choice questions (simple and multiple selection including seven yes/no questions), all of which offered to elaborate the
selection(s) in a short free text comment, and 19 text entry questions. For a visual overview of the variety of measurement
setups used, we asked the survey participants to upload a photo of their chamber system. This part of the survey contained 40
150 questions of different formats, consisting of 20 multiple choice questions of which 7 were yes/no questions and all with the
option to elaborate the response in a short free text comment, five questions asking for multiple short text, 14 free-text questions
and one image file upload. To assess the professional background of the group of participants we asked about their professional
status, the country of their home institute as well as their educational and scientific background. ~~Multiple responses were~~
~~allowed.~~ For an overview of the area of application of chamber CH₄ flux measurements, we included questions on the
participants' research questions and the regions and ecosystem types they usually work in. Questions on the chamber
155 dimensions, the chamber equipment, measurement instruments, as well as photos thereof, together with questions on the
measurement procedure and additional variables monitored showed us the variety of measurement setups~~experimental designs~~
used. Additionally, we asked the participants to describe their approaches for flux calculation, quality control, and uncertainty
estimation of the flux estimates.

Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

2.2. Methods of: Survey part 2 - Visual quality control of a standardized data set

To more directly ~~comparassesse~~ the differences ~~int~~ interpretation of chamber data that leads to the discrepancies in measurement setups, data processing, and QC techniques as identified in the first survey part, we provided a standardized set of chamber measurements for visual QC by the survey participants and extrapolated the responses to a larger, representative data set. This second part of the survey included both qualitative and quantitative responses.

The standardized set of chamber CH_4 fluxes was ~~based on~~ composed of 12 selected chamber measurements from our field campaigns at Siikaneva bog (61°50'N, 24°12'E), Southern Finland, in summer 2021 and summer and fall 2022. The measurements were done ~~with using~~ a manual chamber with a volume of 36 ~~L~~ and, equipped with a cooling system to keep the chamber temperature close to constant, two fans to mix the air inside the chamber, and a small opening for pressure equilibration. For the measurements, the chamber was placed on collars that were permanently installed in the ground. In 2021, the connection between chamber and collar was sealed with a rubber skirt and in 2022 the rim between chamber and collar was filled with water to make the connection air tight. The gas concentrations inside the chamber were ~~measured-recorded~~ with an in-line gas analyzer at a frequency of 1 Hz. Besides chamber measurements showing a linear increase in CH_4 concentration over time, we included examples showing a variety of deviations from the linear increase expected for constant diffusive wetland CH_4 emissions.

~~ForThe survey questions for~~ visual QC of the measurements by the survey participants, we provided ~~included~~ the concentrations of CH_4 over time as well as the simultaneously measured concentrations of CO_2 and H_2O in the chamber, a photo of the chamber, and a description of the measurement setup as well as for each measurement example, information on dominant vegetation and water table depth at the measurement plot, date and time of the measurement, transparent vs. opaque chamber, gas analyzer model and a photo of the measurement plot (Figure A1a). We asked the participants if they would keep the respective measurement for flux calculation or if they would discard it and why they would do so (Figure A1b). If they decided to keep the measurement, we asked them to select the part of the measurement that they would use to calculate the CH_4 flux by submitting the start and end times of this period in seconds after chamber closure and submit this for a calculation of CH_4 flux based on their response as the quantitative portion of the response.

2.3. Statistical analyses

2.3.1 Cleaning of the data set

We anonymized the survey responses by separating the demographic information including the country of the home institute, the scientific background, the highest education level, the time since PhD completion, and the current professional role of the participants from each other and from the rest of the survey ~~responses~~ results. We furthermore removed the question for specific research sites before publishing the data and replaced two names of specific research sites given as a description of the main study regions by terms for a larger region. In one response, we removed the name of another researcher mentioned by one of the participants.

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

We harmonized and/or categorized ~~certain~~some free text responses including the responses on the chamber shape, the chamber area, chamber volume, the closure time of the chamber, and the frequency of the gas concentration measurements inside the chamber. ~~From the~~We used chamber volume and chamber area ~~we to~~ calculate the effective chamber height. We corrected ~~some~~ obvious writing mistakes throughout the survey as part of the standardization. In ~~questions on the visual~~ QC procedures ~~part of the survey~~, we standardized the information on the exclusion of the beginning of the measurement from flux calculation as well as the length of the excluded time period. We also adjusted the responses to questions whether to keep or to discard a measurement in the visual QC ~~exercise~~part when the free text responses clearly revealed that the wrong box ~~had been~~was ticked by mistake. We set the CH₄ flux to zero in two cases where survey participants clearly stated in their free text responses that this is ~~what they would do~~how they would handle the presented measurement.

2.3.2 Evaluating the visual QC exercise

We quantitatively and qualitatively evaluated the responses to the visual QC portion of the survey. We summarized the ~~reasons for keeping or discarding a measurement as elaborated in the~~ free text responses ~~to in~~ the visual QC part ~~for the qualitative responses related to the reasons for keeping or discarding a flux of the survey~~. Then, we numerically evaluated the visual QC performed on the 12 example measurements. This allowed us to quantify the variation in fluxes due to quality control and differences in fitting approaches among researchers. For this, we calculated ~~the~~ CH₄ fluxes for each researcher for each of the 12 ~~selected~~example-QC measurements using the time periods selected by the researcher.

~~To calculate the fluxes, w~~We used a standard linear fitting approach ~~for the flux calculation and~~; accounting ~~ing~~ for differences in temperature and pressure among the measurements (Holland et al., 1999). The ~~I~~ideal Gas ~~L~~aw was used to convert the rate of change in CH₄ concentrations ($\frac{dc_{CH_4}}{dt}$) in ppm s⁻¹ to the molecular CH₄ flux (F_{CH_4}) in mol m⁻² s⁻¹ for each measurement example i ($i=1, \dots, n$, where $n=12$) and each survey participant j ($j=1, \dots, m$, where $m=36$).

$$F_{CH_4 i,j} = \frac{dc_{CH_4}}{dt}_{i,j} \times 10^{-6} \times \frac{p}{R \times T_i} \times \frac{V_i}{A_i},$$

where p represents the standard atmospheric pressure of 101325 Pa, T (degrees K) is the ~~mean~~mean of the temperature inside the chamber during the closure, ~~and~~ A is the surface area of the chamber in m². V_i is the volume of the chamber used in measurement i , calculated by $V_i = A \times h_i$, where h_i is the effective height of the chamber headspace during measurement i (in m), calculated as the mean of the height above the soil surface or vegetation cover that was measured at three points around the chamber for each measurement plot. R is the Ideal Gas Constant of 8.314 kg m² mol⁻¹ K s⁻². We then converted the molecular CH₄ flux to the more commonly used mass flux of CH₄ using the molar mass of CH₄ of 16.04 g mol⁻¹. For each measurement example and each participant, $\frac{dc_{CH_4}}{dt}$ was estimated as the slope of a linear fit (lm function from stats package in R version 4.3.0) to the CH₄ concentrations within the time period selected by the researcher. For reasons of consistency, we used a linear fit even in the 12 cases that a participant suggested to use a nonlinear fit instead (7% of the total of 173 times that start and end

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

times for flux calculation were given by ~~at~~ the participant(s). When a measurement was accepted by an expert but no start and end time was given for flux calculation, we estimated the flux based on the entire chamber measurement.

We used the fluxes calculated from the quantitative responses to assess the ~~uncertainty-variability in CH₄ flux estimates~~ and QC procedures due to different researchers processing the measurement data, that is (1) the ~~uncertainty-variability in flux estimates introduced by~~ due to different researchers (~~4~~) selecting time periods for flux calculations, and 2) ~~the variability in the share of measurements kept for flux calculation during QC-deciding whether to keep or to discard a measurement (quality control)~~. In a representative data set of 788 chamber measurements, collected at Siikaneva bog in 2021 and 2022 (Jentzsch et al., 2024a), we visually identified and categorized the following eight ~~classes~~ classes of measurement ~~scenarios~~ based on the shape of the CH₄ concentrations measured in the chamber headspace over time: “Linear increase”, “Linear decrease”, “Nonlinear increase – decreasing slope”, “Nonlinear increase – increasing slope”, “Initial jump”, “Jump(s)”, “Inconsistent trend”, and “~~Low variation~~ No trend”. During the majority (60%) of measurements in the Siikaneva data set CH₄ concentrations increased linearly over time (Table A1). The second largest group, represented by 18% of the measurements, showed a nonlinear, weakening increase in CH₄ concentrations over the time of the chamber closure. During 8% of the measurements an abrupt jump in CH₄ concentrations in the beginning or one or several jumps at a later time during the measurement were detected, respectively. A nonlinear increase in CH₄ concentrations, that strengthened over time was found in 3% of the measurements and 2% of the measurements had an inconsistent and abruptly changing concentration trend. Low concentration changes, showing no clear trend, and a linear decrease in CH₄ concentrations, together, were represented by less than 1% of the measurements (Table 1). From the Siikaneva data set, we selected 12 measurement examples so that each measurement ~~scenario~~ class was represented at least once in the visual QC exercise (Table A1).

For each measurement ~~scenario~~ class, we estimated the ~~uncertainty-variability in flux estimates~~ introduced by different researchers choosing different time periods within the same measurement for flux calculation using the coefficient of variance (CV) across the fluxes calculated for each survey participant. To extrapolate this ~~variability~~ ~~uncertainty~~ to a representative data set (the presented fluxes were chosen to capture the range of observed behavior, rather than represent the observations as explained above), we calculated the weighted sum of the CVs based on the relative occurrence of each measurement ~~scenario~~ class within the Siikaneva data set (Table A1). To assess the ~~uncertainty-variability indue to QC procedures~~ ~~quality control~~, we extrapolated the percentage of measurements kept for flux calculation to a representative data set for each participant, again using the relative occurrence of each measurement ~~scenario~~ class within the Siikaneva data set. We then calculated the CV between the percentages of measurements kept across all survey participants.

3 Results

A total number of 36 expert researchers participated in the survey. All of them completed the survey parts on demographic information and their field sites for flux measurements. Most participants (35-participants) answered the questions concerning their flux measurement setup, and 30 responded about their flux calculation and QC approach. Participation decreased to 28

Formatted: Subscript

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: Subscript

Formatted: Subscript

Formatted: Subscript

Formatted: Subscript

Formatted: Subscript

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

experts for the visual ~~quality-control~~^{QC} ~~part~~^{exercise} and an additional two ~~participants~~ dropped out after the second example measurement, resulting in a survey completion rate of 72%.

3.1 Results of: Survey part 1 - The survey participants and their chamber measurements

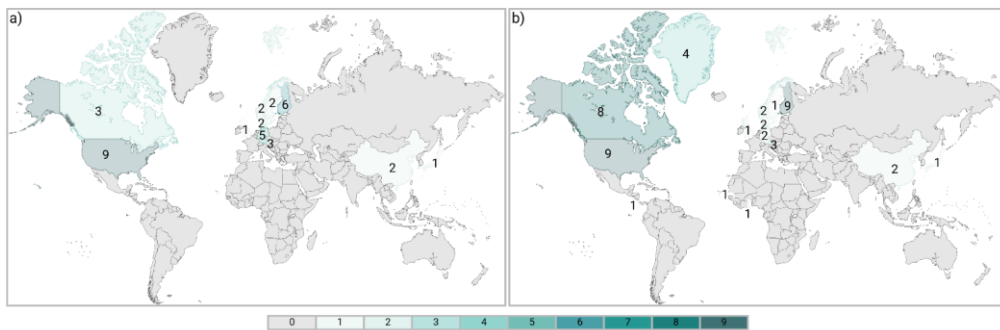
3.1.1 Demography

The ~~survey respondents~~^{participants} work for universities (25 participants), research ~~centers~~^{centers} (11 participants), or companies (one participant) that are located in North America, central and northern Europe, and eastern Asia (Figure 1a). Most (89%) of the participants have a PhD title, 41% of whom completed their PhD within the last seven years, 25% between 7 and 15 year ago, and 34% more than 15 year ago (Figure 2a). Nearly all (94%) of the participants are researchers, two of whom are PhD students (Figure 2b). One participant each specified their current position as Bachelor student, professor, leader in industry, coordinator, and consultant, respectively. With 58%, the majority of the survey participants has a background in Geosciences, followed by ~~b~~^Biology (25%), ~~e~~^Ecology (11%), ~~m~~^Meteorology (8%), ~~e~~^Environmental sciences (6%), and ~~p~~^Physics (6%). One participant each has a background in forestry, biogeosciences, and agricultural sciences. Half of the participants (52%) are part of one or several of the flux networks and databases FluxNet, ICOS, AmeriFlux, OzFlux/TERN, European Fluxes Database Cluster, and LTER.

Formatted: Indent: First line: 0,5 cm

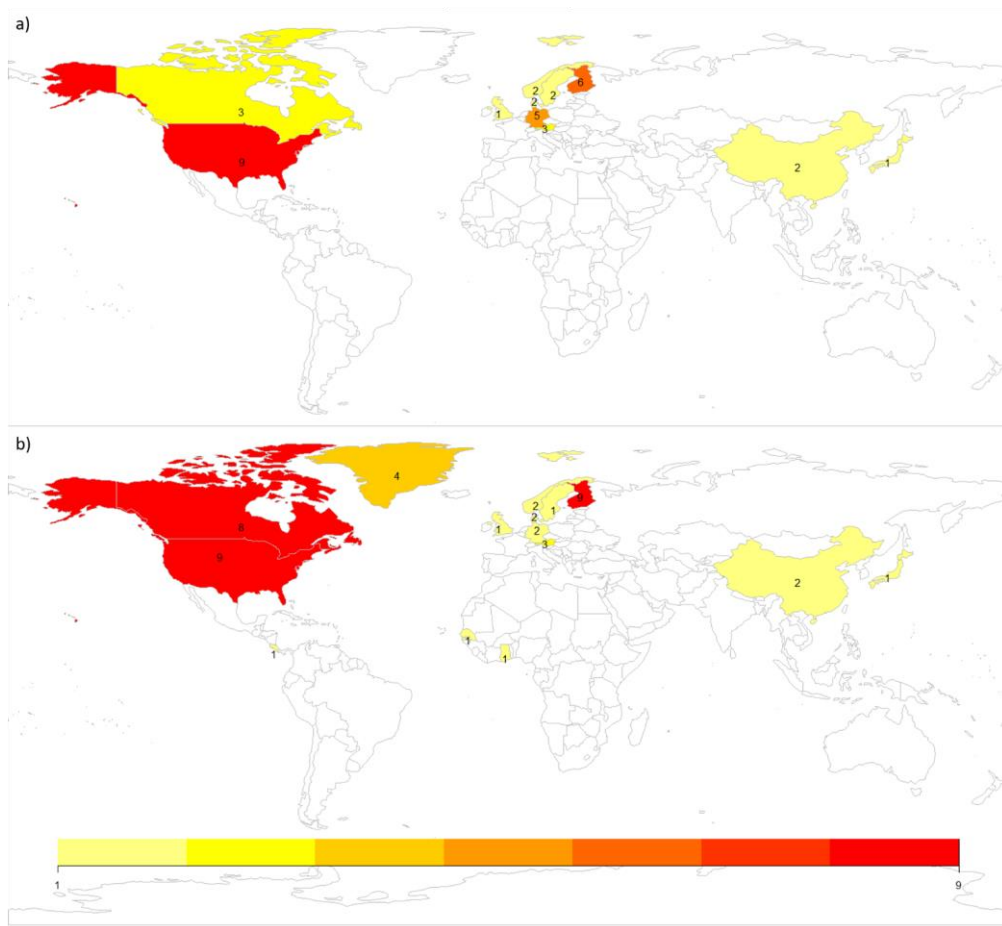
Formatted: English (United States)

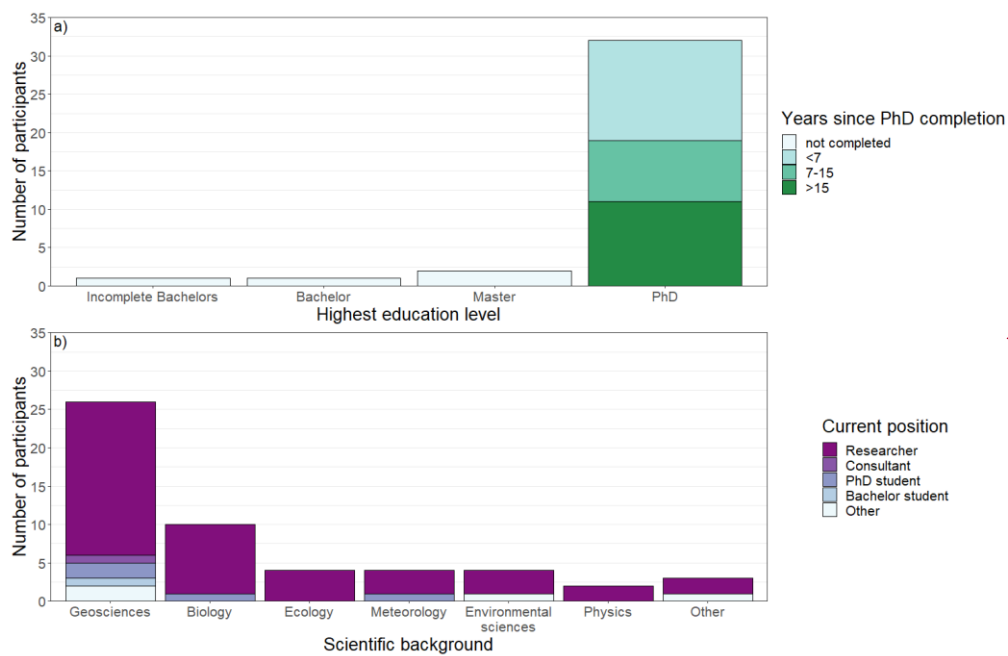
Formatted: Indent: First line: 0,5 cm, Don't keep with ne
Don't keep lines together



Formatted: English (United States)

Formatted: Keep with next





Formatted: English (United States)

Formatted: Keep with next

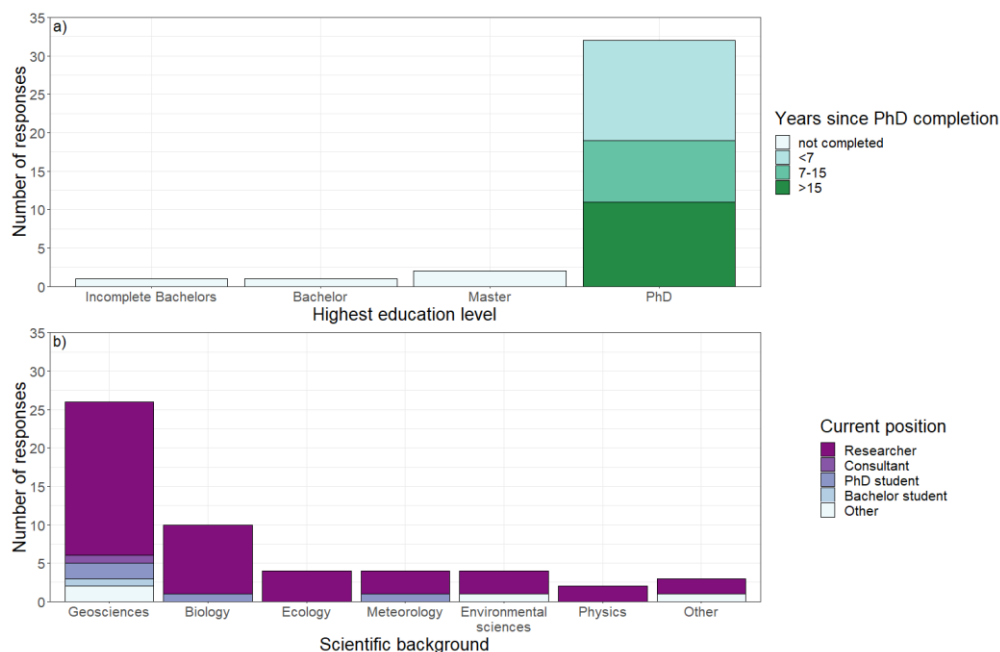
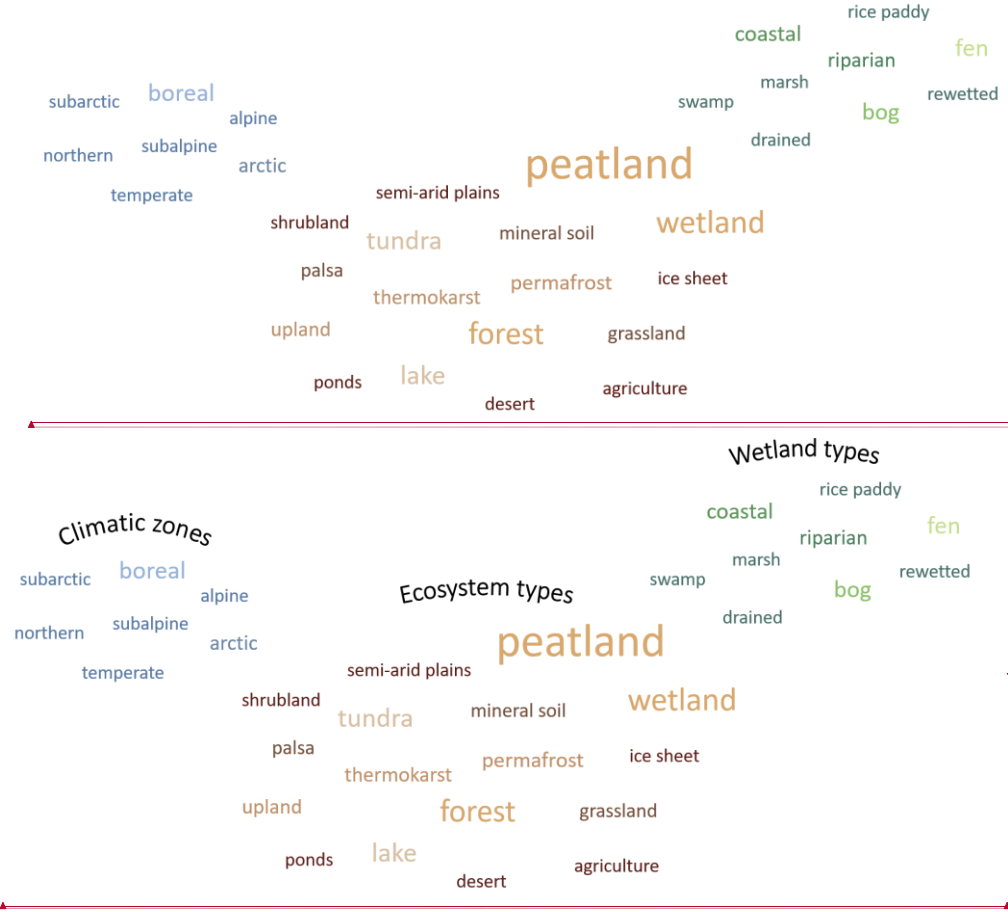


Figure 2: Histograms of the highest education level of the participants, split by the years since their PhD completion (a) and of their scientific background by current position (b). Some participants gave multiple answers regarding their scientific background, causing the total number of responses to exceed the total number of 36 participants.

3.1.2 Flux measurement sites

Most (83%) of the participants do field measurements in the same country of-as their home institute, among them all participants working for institutions in Asia, Canada, Finland, Norway, Denmark, Austria, and the United Kingdom (Figure 1b). Four participants additionally reported field measurements in Greenland and one participant in Ghana, Costa Rica, and Senegal, which were not among the countries of home institutes of the participants. Six participants from the US, Germany, and Sweden had their main research sites in Canada, Finland or Greenland, according to their research questions and ecosystems of interest. The majority (83%) of the participants focus their research on peatlands and wetlands, mainly fens or bogs (50%), and littoral wetlands (31%) (Figure 3). A few (14%) of the participants measured in (semi-)arid regions, upland areas, and at sites with mineral soil instead of or in addition to wetlands. Some (33%) of the participants explicitly mentioned field measurements in permafrost-affected landscapes; similarly, 33% of the participants explicitly mentioned that they measure in “northern”, “boreal”, “arctic”, or “subarctic” regions and 6% measure in “alpine” or “subalpine” terrain. Some

290 (25%) of the participants do aquatic measurements and 19% measure at anthropogenically managed sites such as on agricultural land, in drained and in rewetted peatlands. Specific ecosystems researched by two participants are rice paddies and reed ecosystems.



295 **Figure 3: Word clouds of the study areas with the brownish, middle part of the word cloud representing the climatic zones of the study sites, the studied ecosystem types, the blue part in the top-left corner showing the climatic zones of the study sites, and the green part in the top-right specifying the types of wetlands and peatlands that are researched by the participants.**

3.1.3 Research goals

The overarching research goals that the survey participants address with their flux measurements are to better understand the processes involved in greenhouse gas cycling, to better understand and quantify the effect of changes on greenhouse gas dynamics, to estimate greenhouse gas budgets, and to research the methodology for gas flux measurements. To investigate the environmental and ecological controls on the greenhouse gas exchange is the main goal of 28% of the participants, mainly in peatlands and wetlands and considering environmental conditions, vegetation properties, and the microbial community among others. Specific ecosystems researched by two participants are rice paddies and reed ecosystems. The main aim of 53% of the participants is to understand and/or to quantify the effect of natural and anthropogenically induced change on greenhouse gas dynamics. The changes considered involve climate change, more specifically, warming, vegetation changes, elevated atmospheric CO₂ concentrations, permafrost thaw, and intensifying disturbances, such as wildfires, as well as peatland management, land-use change, and oil and gas exploration. Estimating greenhouse gas budgets is the goal of 22% of the participants but this goal varies differed in spatial and temporal scales from annual budgets of northern ecosystems to budgets of wetlands, microseepage, i.e. diffusive CH₄ fluxes over productive hydrocarbon basins, as an estimate of natural geologic CH₄ emissions, or permafrost and periglacial ecosystems, including thermokarst lakes, thawing permafrost peatlands, and degrading subaqueous permafrost. One participant uses the flux measurements to research methodologies for gas flux measurements, investigating their accuracy, minimum detectable fluxes, curve fitting approaches, as well as engineering challenges around automation and minimizing measurement artefacts.

3.1.4 Flux measurement setup – Guidelines and implementation

There are several guidelines on best practices for chamber measurements that involve recommendations on the chamber setup (e.g. de Klein and Harvey, 2012; Fiedler et al., 2022). The aim of these guidelines is to keep the flux between soil, vegetation and chamber headspace as close as possible to the “real” flux that would be found in the absence of a chamber. This is achieved by minimizing chamber-induced artefacts. Such artefacts include an increasing deviation of environmental conditions inside the chamber from the ambient conditions over the time of the chamber closure and a disturbance of the system during the chamber placement. These chamber effects are reduced by equipping the chamber with additional features such as a vent and shading or active cooling to avoid a pressure and temperature change inside the chamber, respectively, a fan for mixing to avoid the buildup of a stable layering within the chamber (Clough et al., 2020). At the same time, the influence of remaining chamber artefacts can be reduced by a balanced combination of closure time and chamber dimensions and the remaining influence of chamber artefacts can be assessed depending on sampling frequency and additional variables measured. The efficiency of chamber setup recommendations at avoiding chamber artefacts have in part been demonstrated by experimental or modelling studies (e.g., Hutchinson and Livingston, 2001; Pumpanen et al., 2004). Our expert survey revealed that researchers use different instrumental setups, most of them implementing the recommended measures (Figure 4, Figure A2). The participants use different instrumental setups to measure gas fluxes with the chamber technique (for examples see

Formatted: Heading 3

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

figure A2). Of the 33 participants who responded to the question, 82% use manual chambers and 33% use chambers that open and close automatically. The shape of the chamber differs between participants, with 64% using square or rectangular chambers and 55% using cylindrical chambers. The volume of the chambers ranges between 8 and 1800 L with a median of 64 L and an interquartile range (IQR) of 105 L. 93% of the chambers used are smaller than 260 l (Figure A3). The participants equipped their chambers with different devices and use varying approaches to sample the chamber air and measure the concentrations of different greenhouse gases (Figure 4). The majority of the participants (80%) use an in-line gas analyser for on-site, continuous, and high-frequency measurements of the gas concentrations. All but one of them use a closed sample loop which returns the air to the chamber after circulation through the gas analyser. Fewer (26%) of the participants manually sample the chamber air for analysis of a gas chromatograph. One participant uses open-path LI-COR gas analysers installed inside a large chamber. The gas analysers used by the survey participants continuously measure the gas concentrations at high frequencies between five times per second and once every 15 s and therefore require shorter closure times of 0.5 to 12.5 min compared to the closure times of 16 to 50 min used when manually sampling the chamber air every 4 to 10 min (Figure A4). Most survey participants (58%) calibrate their gas analysers once per year and 24% do so once before each measurement campaign. A few (10%) of the participants calibrate the gas analyser less often e.g. when serviced every 1 to 3 years and 12% calibrate more frequently ranging from weekly to daily to calibration after each flux measurement. As required for the survey participation, all participants measure the CH₄ concentrations in the chamber, 97% additionally measure CO₂ and 33% measure the N₂O concentrations. One participant each additionally measures BVOC and Ethane.

Most participants (83%) record the temperature inside the chamber and 29% measure the pressure inside the chamber with measurement frequencies ranging from every second to once per chamber closure.

Pressure vent,

A gas flux into or out of a closed chamber, would slowly alter the air pressure inside the chamber over time as well as more rapidly when the chamber is closed. As such as change in pressure can affect the gas flux between soil and chamber, it is recommended to install a vent, that is a small opening in the chamber, that allows for pressure equilibration but does not allow for significant mixing of ambient air into the chamber to keep the pressure inside the chamber close to ambient air pressure. Clough et al. (2020) recommend the simultaneous use of two types of vents as they tackle different pressure-related chamber artefacts, a larger one that is open only during chamber placement and a smaller one that remains open during the measurement. Vents for pressure equilibration are only used by half of the participants (Figure 4). Half of the participants use a vent to keep the air pressure inside the chamber close to ambient conditions and thereby prevent pressure changes from altering the natural gas flux. Different methods for pressure equilibration employed by the respondents used were a hole in the chamber that is sealed after chamber placement, explicitly mentioned by two participants, and a long line of tubing that is constantly open to the atmosphere allowing for pressure equilibration while preventing that too much ambient air enters the chamber, explicitly mentioned by one participant. The responses indicate that the two types of vents are considered rather as alternatives for vent designs than as two measures that tackle different pressure-related chamber artefacts and that should

- Formatted: Font: Not Bold, English (United States), Not Highlight
- Formatted: Font: Italic, English (United States), Not Highlight
- Formatted: Heading 3
- Formatted: Font: Not Bold, Italic, English (United States), Highlight
- Formatted: English (United States)
- Formatted: Indent: First line: 0,5 cm
- Formatted: English (United States), Not Highlight
- Formatted: English (United States)
- Formatted: English (United States), Not Highlight
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States), Not Highlight
- Formatted: English (United States), Not Highlight
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States), Not Highlight
- Formatted: English (United States)
- Formatted: English (United States), Not Highlight
- Formatted: English (United States), Not Highlight
- Formatted: English (United States)
- Formatted: English (United States), Not Highlight
- Formatted: English (United States)
- Formatted: English (United States), Not Highlight
- Formatted: English (United States)

therefore both be applied simultaneously. One reason for the low implementation rate of pressure vents could be a fear of causing a so-called Venturi effect, where wind passing over the vent outlet can depressurize the chamber, leading to an increased gas flow from the soil into the chamber (Bain et al., 2005; Conen and Smith, 1998). Clear guidelines exist however on how to avoid the Venturi effect by adjusting the vent design (Xu et al., 2006).

Cooling

Especially in summer, when air temperatures are high, a transparent chamber might act as a small greenhouse, causing the temperature inside the chamber to rise and increasingly deviate from the ambient air temperature over the time of the chamber closure, inducing a temperature gradient between the interior and the exterior of the chamber. A change in chamber temperature should be avoided as it can affect the gas flux through influencing processes like plant processes and evaporation or condensation. About one fifth of the survey participants addresses this issue in their chamber setup. As a way to avoid a temperature increase by insulation, 3% of the participants use opaque or reflecting chambers. Some applications, however, require the use of transparent chambers. This is the case for example when determining NEE. Furthermore, blocking out the incoming radiation can potentially reduce active CH₄ transport through plant aerenchyma thereby reducing the measured CH₄ emissions (Clough et al., 2020). 17% of the respondents therefore use active cooling of a non-insulated, transparent chamber. Types of cooling systems mentioned were Peltier elements, circulation of the chamber air through a tank filled with ice-water, and fans circulating the cold air from ice packs placed inside the chamber. However, an active cooling of the chamber air bears the risk of overcompensating for a temperature increase and causing condensation inside the chamber or sampling tubes (Fiedler et al., 2022). It is therefore recommended to use active cooling only if chamber cannot be insulated and/or if long chamber deployment periods are needed (Maier et al., 2022). The effectiveness of insulation or cooling should be evaluated by comparing surface soil temperatures inside and outside the chambers (Clough et al., 2020).

Chamber pressure and temperature measurements

Recording the temperature and the pressure inside the chamber over the time of the chamber closure is essential for correcting for temperature and pressure using the ideal gas law when calculating CH₄ fluxes as well as for detecting remaining changes in pressure and temperature over time that could not be eliminated with a pressure vent and insulation or cooling of the chamber headspace. Most participants record the temperature inside the chamber, while only a little less than one third of them measure the chamber pressure with measurement frequencies ranging from every second to once per chamber closure. While one temperature and pressure measurement during chamber closure might be sufficient for use in the ideal gas law, higher frequency measurements are needed in order to consider the stability in environmental conditions inside the chamber as an indicator of flux quality. Only two participants can therefore account for temperature and/or pressure changes over the time of the chamber closure by individually correcting each concentration measurement as they document the chamber temperature and/or pressure at the same frequency as the gas concentrations. Most notably, almost one fifth of the survey

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Field Code Changed

Formatted

Formatted: Font: Not Bold, Italic, English (United States), Highlight

Formatted: Font: Italic, English (United States), Not Highlight

Formatted: Heading 3

Formatted: English (United States), Not Highlight

Formatted: Indent: First line: 0,5 cm

Formatted

Field Code Changed

Formatted

Field Code Changed

Formatted

Field Code Changed

Formatted

Formatted: Font: Not Bold, Italic, English (United States)

Formatted: Heading 3

Formatted: Font: Italic, English (United States)

Formatted: English (United States), Not Highlight

Formatted: Indent: First line: 0,5 cm

Formatted

395 participants does not measure the chamber temperature at all (Figure 4), which can lead to large uncertainties considering the
strong linear effect of temperature on the flux magnitude through the ideal gas law.

Mixing

400 In the absence of air movement in a closed chamber, a concentration gradient can develop inside the chamber, which might
influence the further gas flux between soil and chamber headspace. A well-mixed headspace is furthermore needed to ensure
that a representative gas sample can be taken. While most researchers use fans to mix the air inside their chamber, some
researchers argued that the air flow from circulation through a closed loop with the gas analyzer was sufficient to mix the
chamber air so that particularly small chambers did not need a fan. This statement highlights that further research is needed to
investigate the strength of turbulence that is adequate for particular chamber dimensions to ensure proper mixing of the
chamber air while preventing that additional gas is artificially released from the soil (Christiansen et al., 2011; Maier et al.,
405 2022). The majority of the participants (80%) uses a fan to mix the air inside their chamber. Three participants argued that the
air flow from circulation through a closed loop with the gas analyser was sufficient to mix the chamber air, particularly in
small chambers. Some (20%) of the participants use a cooling system to prevent the temperature of the air inside the chamber
from rising too much above the ambient air temperature. Types of cooling systems mentioned were Peltier elements, circulation
of the chamber air through a tank filled with ice water, fans circulating the cold air from ice packs placed inside the chamber,
410 and an opaque or reflecting cover on the chamber.

Seal

To reliably quantify the momentary gas exchange between a defined soil surface and the atmosphere, the mixing of chamber
air with ambient air needs to be avoided. To achieve this, it is recommended to insert a chamber base into the ground to restrict
lateral gas transport inside the soil and to additionally ensure an airtight connection between the chamber and its base (Clough
415 et al., 2020). Two thirds of the participants (66%) follow this recommendation and place their chamber on top of a
collar base that they inserted into the ground soil between one hour and one year before the measurement. The more time that
passes between base insertion and measurement the less a potential disturbance of the ground and its concentration gradient
will affect the measurement. The fact that the chamber setups employed by one third of the participants do not involve a collar
or a seal might be less problematic than it appears since many participants measure in wetlands or on open water, and the
420 required insertion depth of the chamber into the soil as well as the necessity of a gastight seal are low under water saturated
conditions and at low soil porosities (Clough et al., 2020). Two thirds of the participants aim to make the connection between
the chamber and the collar or the soil gastight by using one or several types of sealing. Besides gaskets and water seals, a
plastic sheet weighed down by a chain, a stocking filled with sand, and foam in the collar groove were mentioned as sealing
methods. Every chamber setup should be tested for gas tightness before it is deployed in the field, as suggested by Clough et
425 al. (2020). The majority of the participants (63%) aim to make the connection between the chamber and the collar or the soil

Formatted: English (United States), Not Highlight

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Field Code Changed

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

Formatted

gastight by using one or several types of sealing. Besides gaskets and water seals, a plastic sheet weighed down by a chain, a stocking filled with sand, and foam in the collar groove were mentioned as sealing methods.

Chamber dimensions

One challenge in developing chamber measurement protocols is to find a balance between a chamber closure time that is short enough to keep the influence of chamber artefacts low but long enough to reach gas concentrations within the chamber headspace that are above the detection limit of the gas analyzer or gas chromatograph used. One way to reduce the minimum time of chamber closure required to exceed the detection limit of the instrument is through reducing the chamber volume.

The volume of the chambers used by the participants ranged from 8 to 1800 L with a median of 64 L and an interquartile range (IQR) of 105 L. 93% of the chambers used are smaller than 260 L (Figure A3). However, more specific recommendations exist on the chamber dimensions besides requirements on its overall volume: To minimize the error caused by potential leakage and maximize the area sampled, an area/perimeter ratio of ≥ 10 cm is recommended, which equates to a diameter of ≥ 40 cm for a cylindrical chamber. Two thirds of the chambers used by the survey participants respect this recommendation and the majority (75%) of chambers with a smaller-than-recommended area/perimeter ratio are cylindrical. Furthermore, a chamber height to deployment time of ≥ 40 cm h⁻¹ is recommended to maximize the flux detection while minimizing the perturbation of environmental variables. This recommendation is followed in 93% of the measurement setups used by the participants. The two remaining setups had too long closure times considering the relatively flat/low chambers. However, flexibility in chamber dimensions and closure time is often limited by the specific conditions of the research site: The minimum closure time needed depends on the flux magnitude of the gas of interest and on the sensibility/sensitivity of the analyzer and the chamber height has to be chosen to accommodate the vegetation while its area might have to be adapted to the surface structure.

Sampling techniques and chamber closure times

Besides reducing the chamber volume, increasing the measurement frequency of the gas concentrations can reduce the required chamber closure time as in most researched environments CH₄ emissions are high enough so that the minimum detectable flux is reached rather quickly. Much higher sampling frequencies can be achieved through the use of in-line gas analyzers as opposed to manual sampling of the chamber headspace. The majority of the survey participants use an in-line gas analyzer for continuous and on-site measurements of the gas concentrations inside the chamber (Figure 4). All but one of these participants employ a closed sample loop which returns the air to the chamber after circulation through the gas analyzer. One participant uses open-path LI-COR gas analyzers installed inside a large chamber. The gas analyzers used by the respondents record the gas concentrations at frequencies between five times per second and once every 15 s. The chamber measurements therefore use shorter closure times of 0.5 to 12.5 min compared to the closure times of 16 to 50 min used by the fewer participants who manually sampling the chamber air every 4 to 10 min (Figure A4). Two participants using manual sampling keep their chamber closed for more than 40 min which is considered as too long by Rochette and Eriksen-Hamel (2008) while earlier guidelines allowed for up to 1 hour closure time (Holland et al., 1999). To avoid overly long closure times that promote

Formatted: Font: Not Bold, Italic, English (United States)

Formatted: Heading 3

Formatted: Font: Italic, English (United States)

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: Font: 10 pt, Not Bold

Formatted: Font: 10 pt, Not Bold

Formatted: Font: 10 pt, Not Bold

Formatted: Font: 10 pt, Not Bold

Formatted: Font: 10 pt, Not Bold

Formatted: Font: 10 pt, Not Bold

Formatted: Font: 10 pt, Not Bold

Formatted: Font: 10 pt, Not Bold

Formatted: English (United States)

Formatted: Superscript

Formatted: English (United States)

Formatted: Font: Not Bold, Italic

Formatted: Heading 3

Formatted: Font: Not Bold

Formatted: Font: Not Bold, Italic

Formatted: Font: Italic

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

460 chamber effects on the measured fluxes, the minimum required closure time should be determined considering the minimum detectable flux (MDF) based on the sensitivity of the analyzer and the chamber height (Christiansen et al., 2015; Nickerson, 2016).

465 An additional advantage of in-line gas analyzers over manual sampling, besides reducing the relevance of chamber artefacts through shortening closure times, is that the higher temporal resolution of the gas concentration recordings can reveal remaining chamber artefacts. This enhances the possibilities to evaluate the quality of a flux estimate or to exclude measurement periods affected by chamber artefacts at the stage of flux processing. In-line gas analyzers furthermore allow for the use of chambers that open and close automatically. Such automated chambers are used by one third of the survey respondents. While being more cost-intensive than manual chambers, automated chambers allow for continuous measurements at a higher temporal resolution.

470 The precision of the measured gas concentrations might differ between the survey participants as they calibrate their gas analyzers or gas chromatographs at different time intervals: Most respondents (58%) calibrate their instruments once per year and 24% do so once before each measurement campaign. A few (10%) of the participants calibrate the instrument less often e.g. when serviced every 1 to 3 years and 12% calibrate more frequently ranging from weekly to daily to calibration after each flux measurement.

Reducing anthropogenic disturbance

475 The survey participants take various precautionsmeasures to minimize additionalpotential disturbances to their chamber measurements that can be caused by the presence of those who measure and their way to operate the chamber system. For wet, terrestrial sites, 28 participants stand on more stable ground while measuring, either by using permanently or temporarily installed boardwalks or wooden boards or by choosing a drier patch or a rock to stand on. Six participants furthermore mentioned that they make sure not to walk close to the measurement plots by using automated chambers or walking rules supported by warning tape. For aquatic measurements, participants avoid anthropogenic disturbance of the sediment and thus of the gas release by pulling the chamber into its measurement location with a rope or sitting in a boat while measuring. In addition, careful placement of the chamber, training of those who measure, maintenance of collars and sealing, and carefully keeping the vegetation away from the chamber sides were used to minimize disturbances to the chamber measurements.

Ancillary data

485 Recording aAdditional variables alongside the chamber measurements can help explain the observed gas fluxes as well as identify potential disturbances to the measurements. are recorded by many participants alongside the chamber measurements in order to identify potential disturbances or to explain the observed gas fluxes (Figure 5). The variety of variables measured by the survey participants (Figure 5) might indicate that, depending on their background and research questions, scientists consider different variables as important in controlling CH₄ fluxes. Almost all survey participants measure variables to characterize the soil, hydrological, and meteorological conditions, covering most of the ancillary data suggested by Maier et

Field Code Changed

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: Font: Not Bold, Italic, English (United States)

Formatted: Font: Italic, English (United States)

Formatted: Heading 3, Indent: First line: 0 cm

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

Formatted: Font: Not Bold, Italic, English (United States)

Formatted: Heading 3, Indent: First line: 0 cm

Formatted: Font: Italic, English (United States)

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

al. (2022). The potential effects of the vegetation cover were however considered by less than one sixth of the respondents. To characterize the soil and hydrological conditions, variables like C and N content, pH value, reduction-oxidation potentials, soil moisture, soil or water temperature and water table depth are measured by 97% of the participants. Meteorological conditions are documented by 88% of the participants recording variables like atmospheric temperature, atmospheric pressure, relative humidity, vapor pressure deficit, rainfall, wind direction, wind speed and photosynthetically active radiation, in particular when partitioning the measured CO₂ fluxes into gross primary productivity and ecosystem respiration. The vegetation cover of the measurement plots is characterized by 15% of the participants through assessing the plant species composition, measuring plant height, specific leaf area, or leaf dry matter content or estimating the leaf area index of the individual plant species or the moss cover by species. In cold regions or seasons, thaw depth, active layer depth, snow depth and/or frost depth are recorded by 9% of the participants. Numerous attempts have been made towards standardizing flux measurements using static chambers by compiling and publishing guidelines on best measurement practices on chamber measurements (e.g. de Klein and Harvey, 2012; Fiedler et al., 2022). Our expert survey revealed that despite the existing guidelines, not all researchers are implementing the recommended measures for various reasons (Figure 4, Table 3). Reasons for not implementing certain chamber equipment or processing advice could be that some researchers might simply be unaware of the recent guidelines on chamber measurements due to a lack of collaboration and a lack of suitable networking platforms. Only half of the survey participants are part of a flux monitoring network, such as FluxNet or AmeriFlux, most of which strongly focus on eddy covariance measurements. Other potential reasons for not implementing recommended measures include that existing equipment might not be suitable for adjustments, e.g. specific chamber sizes or shapes can make some measures inappropriate or impractical, financial constraints might not allow for improvements or new equipment, or there might be site-specific requirements on the chamber design. The various scientific backgrounds of the researchers doing chamber measurements (Figure 2b) might further be contributing to the variety of measurement setups, calculation and QC procedures found in this survey as the educational training likely influences which aspects of the flux measurements, i.e. which chamber artefacts and environmental controls on CH₄ fluxes, are considered most important.

Some shortcomings in measurement setups can likely be related to an uncertainty around the effect of certain chamber equipment on the measured CH₄ flux, leading to concerns that overcompensating for certain chamber artefacts can introduce new sources of error. While most researchers use fans to mix the air inside their chamber, some researchers mentioned that their chambers are small enough to not need a fan. This statement highlights that further research is needed to investigate the strength of turbulence that is adequate for particular chamber dimensions to ensure proper mixing of the chamber air while preventing that additional gas is artificially released from the soil (Christiansen et al., 2011; Maier et al., 2022). Vents for pressure equilibration are likely only used by half of the participants for the similar fear of causing a Venturi effect when wind passes over the vent outlet (Bain et al., 2005; Conen and Smith, 1998). Clear guidelines exist however on how to avoid the Venturi effect by adjusting the vent design (Xu et al., 2006). Furthermore, the two types of vents – the one that is open only during chamber placement and the one that remains open during the measurement – seem to be considered rather as alternatives for vent designs than as two measures that tackle different pressure-related chamber artefacts and that should therefore both

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

525 be applied simultaneously. The survey participants generally avoid the danger of overcompensating for a temperature increase inside the chamber and causing condensation (Fiedler et al., 2022) by using active cooling only in cases where it is required, e.g. because transparent chambers are used. Depending on the environmental conditions, opaque chambers could be used more often for insulation but blocking out the incoming radiation could reduce active CH₄ transport through plant aerenchyma thereby reducing the measured CH₄ emissions (Clough et al., 2020).

530 The ability to detect or correct for any remaining temperature and/or pressure differences between chamber and ambient air varies among the participants as not all of them record the temperature and/or pressure inside their chamber. Only two participants can account for temperature and/or pressure changes over the time of the chamber closure by individually correcting each concentration measurement as they document the chamber temperature and/or pressure at the same frequency as the gas concentrations. Changing temperature and pressure conditions inside the chamber might go unnoticed when using only one temperature and pressure reading to correct the final flux estimate. Most notably, 17% of the survey participants do not measure the chamber temperature at all (Figure 4), which can lead to large uncertainties considering the large linear effect of temperature on the flux magnitude through the ideal gas law.

540 Similar to controlling the temperature inside the chamber the requirements for chamber dimensions, chamber insertion depths into the soil, and chamber seals strongly depend on the environmental conditions of the research site. Chamber dimensions need to be adapted to the surface structure as well as to the vegetation height while allowing for flux detection within reasonable deployment times. As the required insertion depth of the chamber into the soil as well as the necessity of a gastight seal are low under water saturated conditions and at low soil porosities, the fact that one third of the survey participants did not use a collar or a seal on their chamber might be less problematic than it appears since many participants measure in wetlands or on open water. The chamber setup should nevertheless be tested for gas tightness before it is deployed in the field.

545 An increasing use of inline gas analyzers can loosen the requirements on chamber dimensions for CH₄ flux measurements due to shorter deployment times. In most researched environments CH₄ emissions are high enough so that the minimum detectable flux is reached quickly after the chamber closure. The high measurement frequency of inline gas analyzers therefore allows to significantly reduce the chamber deployment time, thereby decreasing the potential effect of chamber artefacts on the CH₄ flux estimates. Chambers can therefore be smaller as shorter measurement times reduce the potential effects of leakage, lateral diffusion, temperature and pressure changes on the flux estimates. Currently, the majority of the survey participants (80%) use inline gas analyzers. Of the 26% of participants who manually sample the chamber air, two participants (22%) keep their chamber closed for more than 40 min which is considered as too long by Rochette and Eriksen Hamel (2008) but earlier guidelines allowed for up to 1 hour closure time (Holland et al., 1999). In addition to increasing the relevance of chamber artefacts due to longer closure times, manual sampling can obscure the influence of chamber artefacts through the lower temporal resolution at which the gas concentrations inside the chamber are monitored. This limits the possibilities to still exclude measurement periods affected by chamber artefacts at the stage of flux processing and quality control.

555 The different variables measured alongside the fluxes might indicate that depending on their background and research questions the survey participants consider different variables as important in controlling CH₄ fluxes. The ancillary variables

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

560

565

also determine which additional information is available to the researchers to evaluate the quality of the CH₄ flux measurements. Almost all survey participants measured variables to characterize the soil, hydrological and meteorological conditions, covering most of the ancillary data suggested by Maier et al. (2022). The potential effects of the vegetation cover were however considered by less than one sixth of the participants only.

Formatted: English (United States)

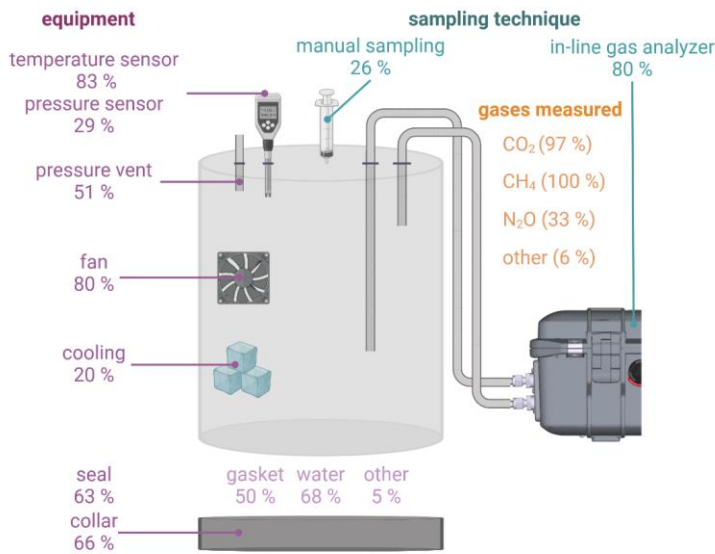
Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)



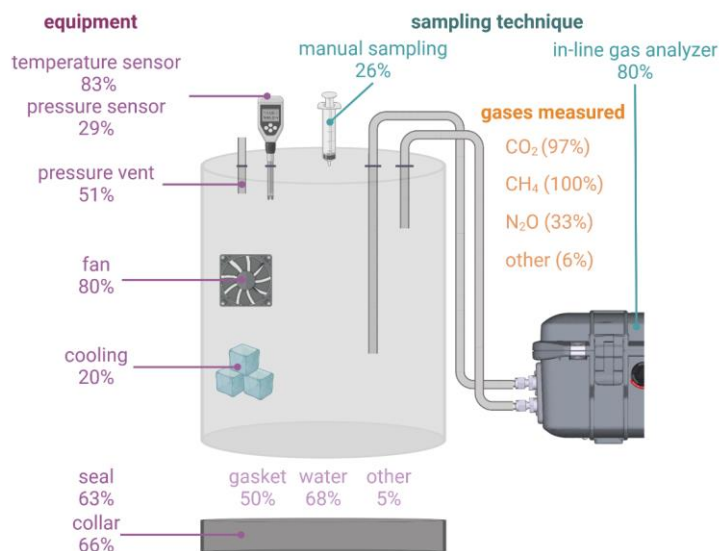


Figure 4: Schematic chamber set-up including the percentage of survey participants using certain types of chamber equipment and sampling approaches for the gas concentrations and measuring different greenhouse gases. Some participants use both manual sampling and in-line gas analyzer measurements at different applications, research sites or measurement campaigns, causing the total share of measurement methods used to exceed 100%. Other gases besides CO₂, CH₄ and N₂O measured by one survey participant each are ethane and BVOC. This figure was created in BioRender.

Formatted: English (United States)

Formatted: Keep with next

Formatted: English (United States)

Formatted: Keep lines together

Formatted: English (United States)

Formatted: English (United States), Subscript

Formatted: English (United States)

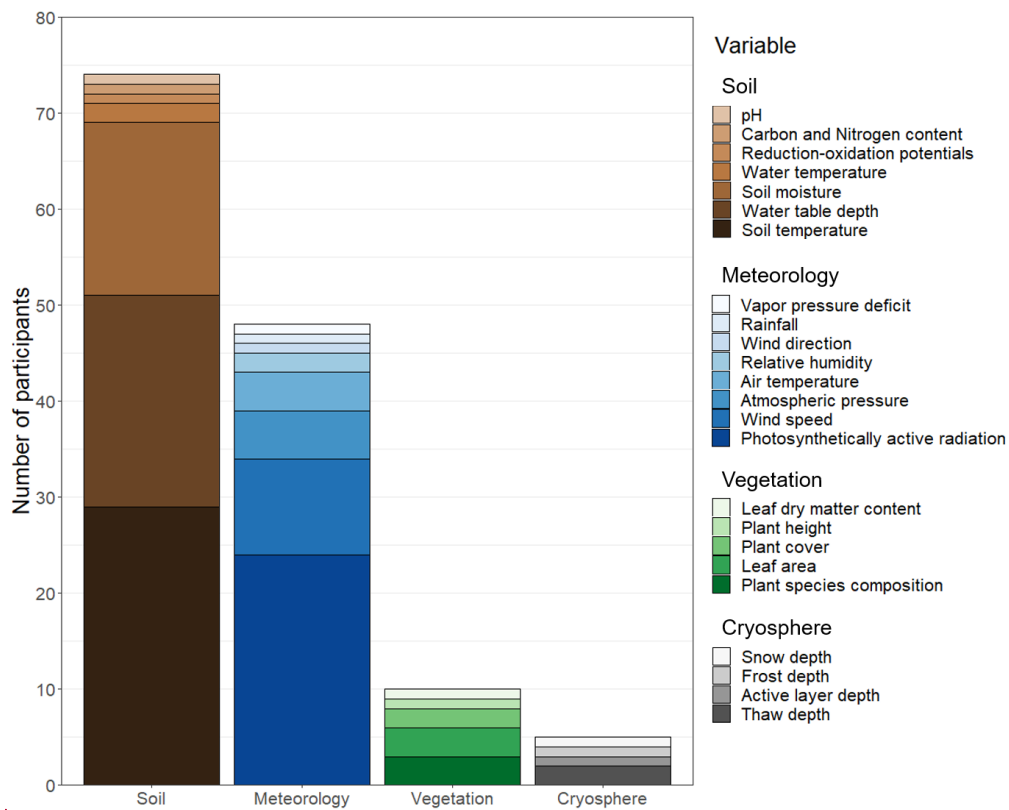
Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: English (United States)



Formatted: English (United States)

Formatted: Keep with next

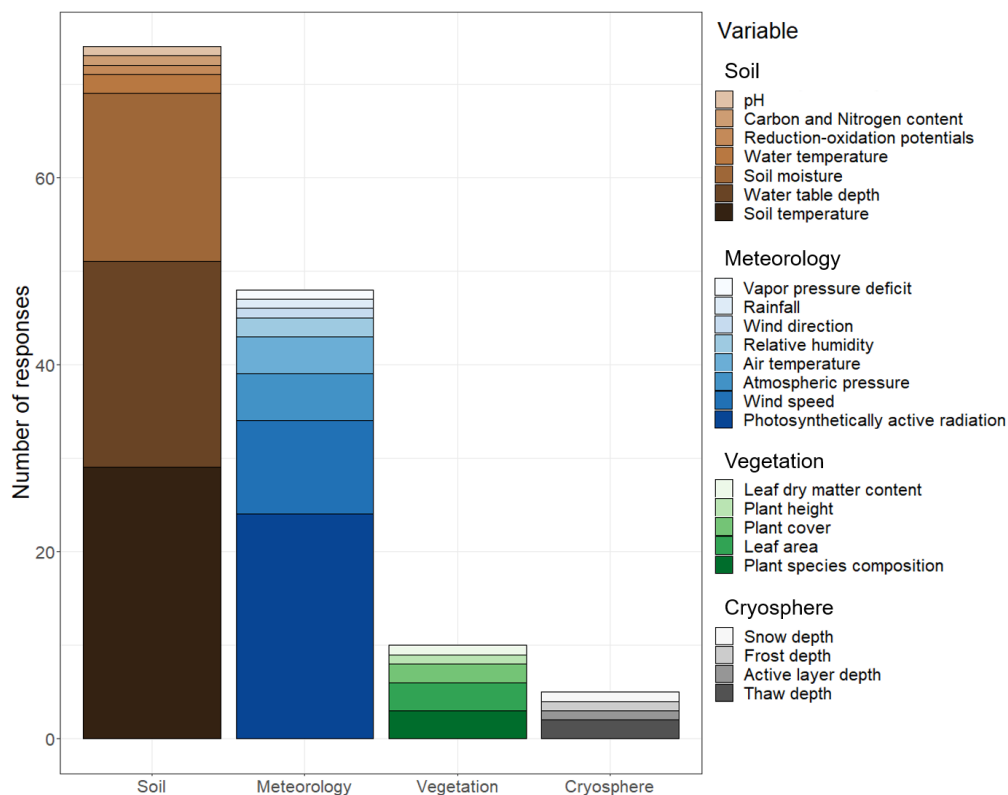


Figure 5: Ancillary data recorded alongside the gas fluxes. Participants were permitted to select multiple variables, allowing the number of responses within one category of ancillary data to exceed the total number of 36 survey participants.

3.1.54 Flux calculation and QC approaches

The qualitative responses on calculation approaches for CH₄ fluxes revealed that slight differences in the flux processing and QC procedures which measurement routines and processing techniques that might result in considerable variation differences in between the CH₄ fluxes derived by different researchers from the same chamber measurement by among different researchers. Gas fluxes are generally estimated from chamber measurements as the slope of the change in gas concentration over the time of the chamber closure and accounting for the water vapor concentrations, the temperature, and the pressure inside the chamber as well as for the chamber dimensions. This approach was modified by the survey participants

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

mainly through selecting a time period of each chamber measurement for flux calculation, choosing a fit function to estimate the change in concentration over time, and determining the accuracy of the temperature and pressure correction by selecting a measurement frequency for the two variables or deciding to use standard values instead (Figure 6). The majority of the participants (90%) use self-written scripts and functions for their flux calculation while 20% of the participants at least partly use existing and published R or Matlab scripts.

Selecting a time period within a chamber measurement for flux calculation, for many respondents, involves discarding the beginning of each measurement to exclude/avoid initial disturbances caused by the chamber placement. a large share of the participants (43%) excludes the beginning of each chamber measurement from their flux calculation ranging from the first 5 s to the first 170 s of the measurement with a median of discarding the first 30 s of each measurement. MostThe majority of the participants (70%) use a linear fit to estimate relationship of the change in gas concentration over the time offrom each chamber measurement. Most remaining respondents Some (23%) of the participants usecompute both a linear fit as well as the initial slope of an exponential fit, either deciding for one based on the goodness of the fit or using the range-difference betweenof the two slopes as an uncertainty estimate for the final flux value. One participant each uses an exponential fit on all chamber measurements, considers the total change in gas concentration as the difference between the gas concentrations at the start and at the end of the chamber closure, or averages multiple linear fits on a one-minute window moving over the measurement at steps of 10 s.

In the step of correcting the measured gas concentrations for the temperature and pressure inside the chamber, most participants use one temperature value per chamber closure, that is either measured during one point of the chamber measurement or derived as the average of several temperature recordings over the time of the chamber closure. As less participants measure the pressure compared to the temperature inside the chamber, more have to rely on ambient pressure recordings or assume standard atmospheric pressure.

Of the 17% of participants who do not measure the temperature inside their chamber, one participant explicitly mentioned that they assume standard temperature in their flux calculations, while two participants use air temperature measurements. The two thirds of survey participants who do not measure the air pressure inside their chamber similarly assume standard pressure or a constant value adapted to their measurement site or use ambient pressure measurements instead. At least 43% of the participants use one temperature value, either measured at the beginning or at the end of the chamber closure, or averaged over the whole chamber measurement for each flux calculation. On the contrary, tAs opposed to assuming constant conditions over the time of the chamber closure, two participants explicitly stated that they individually correct each gas concentration measurement individually for the chamber temperature and/or pressure measured at or interpolated to the same frequency as the concentration measurements.

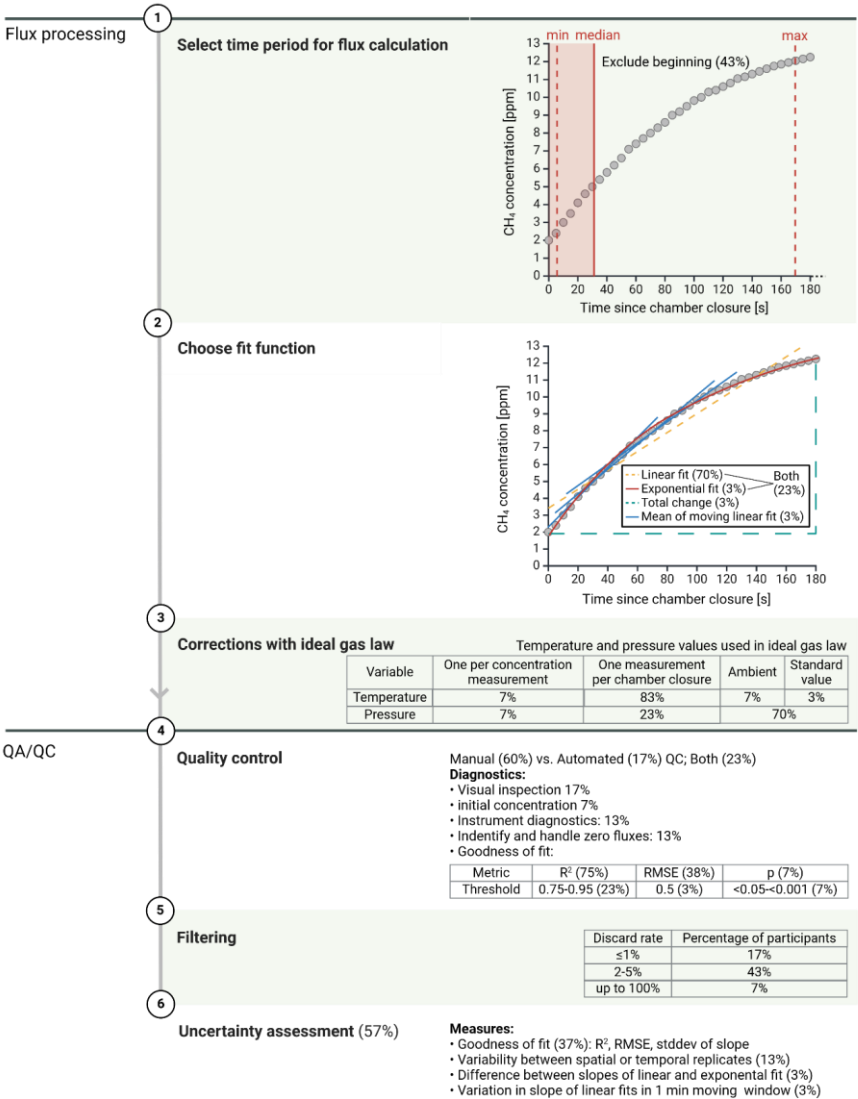
The majority of the participants (90%) use self-written scripts and functions for their flux calculation while 20% of the participants at least partly use existing and published R or Matlab scripts.

Various approaches for QC ofQC of the flux estimate time series of gas concentrations recorded during the chamber measurements were mentioned by the participants. MMost participants (60%) manually check each of their chamber

620 measurements, while ~~others 17% of the participants~~ use an automated QC procedure and ~~some 23%~~ used a combination of both manual and automated diagnosis. ~~Most A majority (53%) of the participants use measures of the goodness of fit to evaluate the quality of their flux estimates, some of whom consider fixed cut-off values of these metrics that decide between keeping or discarding a flux measurement, with the R² and RMSE (root-mean-square error) being the preferred quality measures used by 75% and 38% of the participants, respectively. Fixed cut-off values given by 7 participants for the R² ranged between 0.75 and 0.95 and a RMSE threshold value of 0.5 was given by one participant. Two participants furthermore only accept measurements with a significant change in the CH₄ over time with p values smaller than 0.05 and 0.001, respectively. In addition, 17% of the participants visually inspect the time series of CH₄ concentrations resulting from each chamber measurement, 7% check that the starting concentration in the chamber does not deviate too much from ambient conditions, and 13% each consider instrument diagnostics for their quality assessment or identify cases of low or no CH₄ flux for special treatment. Apart from two participants, the respondents typically discard up to 5% of their data. Most participants (43%) discard between 2 and 5% of their data based on their QC procedure. For 17% of the participants, their QC protocol leads to an exclusion of up to 1% of their measurements while two participants discard up to 100% of their data (Figure A5).~~

630 ~~The uncertainty of each individual flux estimate is assessed by One third of the 57% of the respondents participants assesses the uncertainty of each individual flux estimate, most of them (80%) using metrics for the goodness of fit or the variability between spatial or temporal measurement replicates, based on the uncertainty of the fit parameters expressed as R², RMSE, or standard deviation of the slope. One and the two remaining participants each uses based on the difference between the slopes derived from a linear compared to an exponential fit and/or the variation in several one-minute linear fits in a moving time window as an uncertainty estimate, respectively.~~

- Formatted: Font color: Auto, English (United States)
- Formatted: English (United States)
- Formatted: Font color: Auto, English (United States)
- Formatted: English (United States)
- Formatted: Font color: Auto, English (United States)
- Formatted: English (United States)



Formatted: English (United States)

Formatted: Keep with next

Formatted: Font: 9 pt, Bold

Formatted: Left, Indent: First line: 0 cm, Keep lines together

Formatted: Font: 9 pt, Bold

Formatted: English (United States)

640 **Figure 6: Differences in the workflows used for flux processing, quality control (QC), and quality assurance (QA) by the survey participants. This figure was created in BioRender.**

3.2 Results of: Survey part 2 - Visual quality control of a standardized data set

3.2.1 Explanations for and resulting handling of different measurement scenariosGeneral trends

The visual QC exercise revealed how the measurement examples were handled by the survey participants, their explanations of nonlinear behaviour, as well as how much the flux estimates were affected by the time period chosen for flux calculation depending on the behaviour of the CH₄ concentrations (Table 1).

The visual QC exercise revealed that the handling of the measurement examples (decision to keep or discard a measurement and choice of time period for flux calculation) differed between the survey participants depending on their interpretation of the CH₄ concentration change in the chamber headspace over time (Table 1). Depending on the shape of the concentration curve (linear or nonlinear), the choice of the time period used for flux calculation furthermore had a strong impact on the magnitude and in one case even on the direction of the estimated CH₄ flux (Figure 7, Table A2). Detailed descriptions of the individual measurement examples and their handling by the survey participants can be found in Text A1.

Linear fluxes: Emission and uptake

The majority of the participants (91%) decided to keep the measurements that showed a linear increase in CH₄ concentrations for flux calculation. Due to the linear behaviour, these flux estimates were least affected by the time period that was chosen for the linear fit, showing in a low variation of 2.5% between the flux estimates (Figure 6).

The latter also applied to measurement example showing a linear decrease in CH₄ concentrations over time. However, more participants decided to discard the entire measurement because they did not expect to find net uptake of CH₄ at a wetland site. The free text responses revealed that the conditions, and in particular the water table depths, under which net uptake of CH₄ can occur were debated among the participants.

Nonlinear increase - decreasing slope

Although the flux estimates from measurements that showed a consistent but nonlinear trend in CH₄ concentrations were strongly affected by the time period chosen for flux calculation, Most participants (79%) also kept these measurement examples that showed a consistent but nonlinear and weakening increase in CH₄ concentrations over time. Other than for a linear change in CH₄ concentrationsHere, the magnitude of fluxes estimated from the nonlinear concentration change strongly depended on the time period selected for the flux calculation. The selection of the time period in turn was influenced by how the participants explained the observed nonlinearity. There were two main reasonings among the participants for their choice of the time period with opposing effects on the flux magnitude: (1) About two thirds of explanations for the nonlinear behavior, assumed that the increase in CH₄ concentrations was weakened by either CH₄ saturation of the chamber headspace or leakage of air from the chamber towards the end of the measurement. The participants concluded that this laterlatter part of the measurement was disturbed and should therefore be excluded from the flux calculation, which resulted in higher flux estimates. (2) Conversely, the remaining third of explanations assumed that the stronger increase in CH₄ concentrations at the

Formatted: English (United States)
Formatted: Heading 2
Formatted: Indent: First line: 0,5 cm
Formatted: English (United States), Subscript
Formatted: English (United States)
Formatted: English (United States)
Formatted: English (United States), Subscript
Formatted: English (United States)
Formatted: Font: Not Bold, Italic, English (United States)
Formatted: Heading 3
Formatted: Font: Not Bold, English (United States)
Formatted: Font: Not Bold, Italic, English (United States)
Formatted: Font: Not Bold, English (United States)
Formatted: Font: Not Bold, Italic, English (United States)
Formatted: Font: Italic, English (United States)
Formatted: English (United States)
Formatted: English (United States)
Formatted: Indent: First line: 0,5 cm
Formatted: Subscript
Formatted: English (United States)
Formatted: English (United States)
Formatted: English (United States)
Formatted: Font: Not Bold, English (United States)
Formatted: Heading 3, Indent: First line: 0 cm
Formatted: English (United States)
Formatted: Font: Italic, English (United States)
Formatted: English (United States)
Formatted: Indent: First line: 0,5 cm
Formatted: English (United States), Subscript
Formatted: English (United States)
Formatted: English (United States), Subscript
Formatted: English (United States)
Formatted: English (United States)
Formatted: English (United States), Subscript
Formatted: English (United States)
Formatted: English (United States), Subscript
Formatted: English (United States)
Formatted: English (United States), Subscript
Formatted: English (United States)

beginning of the measurement was caused by an initial disturbance such as ebullition, triggered by the chamber placement. A consequent exclusion of the strong initial increase in CH₄ concentrations measurement start from flux calculation resulted in lower flux estimates as the lower slope during the latter part of the measurement was preferentially selected.

675 *Nonlinear increase - increasing slope*

The survey participants were similarly divided on the appropriate handling of chamber measurements that show a nonlinear increase in CH₄ concentrations but with an increasing slope over time. Accordingly, half of the participants who discarded this measurement argued that they cannot justify choosing a time period for flux calculation as they cannot explain the observed shape in CH₄ concentrations and, considering the nonlinearity, an unsubstantiated selection of a time period could strongly bias the flux estimate. For those who The flux estimates computed for the 65% of participants who kept the measurement and gave start and end times for flux calculation (65% of participants), indeed differed significantly depending on the time period chosen significantly affected the flux estimate. This range between higher and lower flux estimates again resulted from contrasting explanations of the nonlinear concentration change: Higher flux estimates originated from explanations assuming an initial period of adjustment and disturbance caused by the chamber placement, through exclusion of the initial, lower slope in CH₄ concentrations. On the other hand, explanations involving chamber effects on CH₄ cycling processes through alteration of environmental conditions or interference of CH₄ measurements with high H₂O concentrations led to lower flux estimates due to exclusion of the stronger increase in CH₄ concentrations towards the end of the measurement.

685 *Jumps*

The majority of the respondents (65%, 88%, 92%) interpreted the jumps showing in three of the measurement examples as episodic events of ebullitive CH₄ emission while one participant suggested a malfunctioning of the gas analyzer. The survey responses revealed uncertainty around the question under which water table conditions CH₄ ebullition is most likely to occur, indicating a fundamentally different understanding of the causes of ebullition events among the participants. There were two major considerations concerning CH₄ ebullition during chamber measurements: First, the survey participants disagreed on whether ebullition events should be included in flux estimates from chamber measurements or if diffusive and ebullitive flux should be quantified separately, either by isolating periods of ebullitive and diffusive flux in one concentration time series or by separately measuring ebullition, for example using bubble traps. When accounting for both diffusive and ebullitive CH₄ emission by using a linear fit over an entire measurement containing ebullition events, as suggested by 4 to 8% of the participants, flux estimates were up to five times as high as the ones considering the diffusive flux only. Second, the respondents disagreed on whether the remaining part of a measurement after an ebullition event could still be used to quantify the diffusive flux. More than half of the survey participants (54%) kept the linear part of a measurement after an initial ebullition event for flux calculation while 38% of the participants discarded the entire measurement. The latter assumed that the high CH₄ concentrations in the chamber following the ebullition event would decrease the concentration gradient and thus reduce the CH₄ flux between soil and chamber headspace for the rest of the measurement. This decision also influenced the

Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: Font: Not Bold, Italic, English (United States)

Formatted: Heading 3

Formatted: Font: Not Bold, English (United States)

Formatted: Font: Not Bold, Italic, English (United States)

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: Subscript

Formatted: Subscript

Formatted: Subscript

Formatted: Subscript

Formatted: Subscript

Formatted: English (United States)

Formatted: Font: Not Bold, Italic, English (United States)

Formatted: Heading 3

Formatted: Font: Italic, English (United States)

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: Subscript

Formatted: English (United States)

Formatted: English (United States)

740

745

750

755

of a measurement was mentioned several times as an explanation for nonlinear behavior in CH₄ concentrations. Linear changes in CO₂ and H₂O on the other hand were considered a proof the air tightness of the system. To avoid any initial disturbance caused by the chamber placement from influencing the flux estimate, almost half of the survey participants by default excluded the beginning of each measurement (30 ± 85 s (median \pm IQR)xx seconds) from their flux calculations.

Effect of different flux calculations on an example flux dataset

Using the prevalence of different measurement scenarios in the Siikaneva data set (Table A1), we estimated an overall variability in the calculated overall CH₄ fluxes due to difference in time periods used for fitting as well as an overall variability in the inclusion/exclusion of measurements fluxes. Different researchers chose different parts of the same measurement for flux calculation (Figure A1.1 – A1.12), which resulted in an overall flux difference of 17% across the Siikaneva data set (Table A2). The differences in the variability of Siikaneva fluxes due to the variation in the percentages of measurements in the Siikaneva data set passing the visual QC was 28% (Table 1, A2). Therefore, the visual QC and decision to include or exclude fluxes (Table 1) had a larger effect on the overall flux calculation than the difference in time periods selected for fitting. These estimated variability introduced by the selection of different time periods for flux calculations compares with the mean natural temporal variability of 19% but isare lower than the mean natural spatial variability of 88% calculated from automated chamber measurements of CH₄ fluxes in five temperate and Arctic peatlands by Pirk et al. (2016). Pirk et al. (2016) similarly found that both natural spatial and temporal variability in CH₄ fluxes exceed the difference between fluxes estimated using different fit functions. However, it must be noted that the uncertainty estimates derived in our study consider only the effect of differences in visual QC and do not account for different measurement setups or different fit functions used for flux calculation, both of which can add variability to fluxes (e.g., Pihlatie et al., 2013)(e.g Pihlatie et al., 2013).

- Formatted: English (United States)
- Formatted: English (United States), Subscript
- Formatted: English (United States)
- Formatted: English (United States), Subscript
- Formatted: English (United States)
- Formatted: English (United States), Subscript
- Formatted: English (United States)
- Formatted: Highlight
- Formatted: Indent: First line: 0 cm
- Formatted: Font: Italic
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: Subscript
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Field Code Changed
- Formatted: English (United States)
- Field Code Changed
- Formatted: English (United States)
- Field Code Changed
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: Justified, Indent: First line: 0,5 cm, Line space: 1,5 lines

Table 1: Explanations of the overall shape in CH₄ concentrations during chamber closure, reasons to discard, and reasons and approaches to keep measurements as given by the participants as well as percentage of respondents that kept measurements and coefficient of variance (CV) of the flux estimates derived from the survey responses by measurement scenario (For details see Table A2).

Measurement scenario	ID	Explanations	Reasons to discard	Reasons to keep & approach	Kept [%]	CV
Linear increase	VOC1	• Net CH ₄ production & diffusive emission	• Small nonlinearities due to ebullition or saturation	• Consistent linear increase in CH ₄ concentrations	91	2.5
	VOC2		• High initial CH ₄ concentrations	• No indications of significant disturbances or malfunctioning of the instruments		
			• No shading	• Close-to-ambient initial CH ₄ concentrations		
Nonlinear increase – decreasing slope	VOC4	• Saturation	• Saturation	• No clear disturbance	79	44
	VOC5	• Initial disturbance	• No steady state reached	• Nonlinear fit		
	VOC9	• Leakage	• Initial disturbance	• Use more linear part in the beginning		
		• Changing environmental conditions	• Leakage	• Use more linear part at the end		
		• Unsure	• Changing environmental conditions			
			• Unclear which part of the measurement represents real flux			
Initial jump	VOC7	• Ebullition caused by chamber placement	• High CH ₄ concentrations affect concentration gradient	• Use linear part after the jump	62	5
		• Malfunctioning of gas analyzer				
Jump(s)	VOC8	• Ebullition caused by	• High CH ₄ concentrations affect concentration	• Use measurement before first jump	40	42.5
	VOC12	• (anthropogenic) disturbance	• gradient	• Use linear part after the jump(s)		
		• Malfunctioning of gas analyzer	• Ebullition affects pressure inside the chamber	• Use longest linear part in between jumps		
Nonlinear increase – increasing slope	VOC10	• Initial period of mixing or adjusting	• Shape of curve unexpected and strong curvature	• Nonlinear fit	76	19
		• Increase in chamber temperature over time	• makes flux estimate depend strongly on selected time period	• Use more linear part in the beginning		
		• Disturbance of measurement plot/ concentration gradient during chamber placement		• Use more linear part at the end		
		• Chamber affects plant-mediated CH ₄ transport				
		• Leakage				
Inconsistent trend	VOC11	• Net CH ₄ consumption	• No consistent trend of sufficient length	• Keep increasing initial part of the measurement	29	138

		<ul style="list-style-type: none"> • Net CH₄ uptake unexpected → Measurement issue: • Gas analyzer issue • Condensation • Leakage • Malfunctioning fan • Initial disturbance such as ebullition caused by chamber placement 	<ul style="list-style-type: none"> • Unclear which part to use for flux calculation because reason for pattern / timing of disturbance unclear 	<ul style="list-style-type: none"> • Keep later, decreasing part of the measurement 		
Linear decrease	VQC3	<ul style="list-style-type: none"> • Net CH₄ uptake • Leakage • High initial CH₄ concentrations 	<ul style="list-style-type: none"> • Net CH₄ uptake unexpected in wetland • Initial ebullition • Anthropogenic disturbance • Leakage 	<ul style="list-style-type: none"> • Net CH₄ uptake possible 	50	17
Low variation	VQC6	<ul style="list-style-type: none"> • Production and oxidation balance • “zero flux” (uncertainty > flux) • Leakage 	<ul style="list-style-type: none"> • Leakage • Changing trend in CH₄ concentrations 	<ul style="list-style-type: none"> • Manually set flux to zero • Small but real flux exceeding instrument precision 	38	57

Formatted

Formatted: Indent: Left: 0 cm

Formatted: Font: 8 pt

Formatted: List Paragraph, Indent: Left: 0 cm, Hanging: 0,2 cm, Bulleted + Level: 1 + Aligned at: 0 cm + Indent at: 0,63 cm

Formatted: Font: 8 pt

Formatted: List Paragraph, Indent: Left: 0 cm, Hanging: 0,2 cm, Bulleted + Level: 1 + Aligned at: 0 cm + Indent at: 0,63 cm

Formatted: Font: 8 pt

Formatted: Left

Formatted: List Paragraph, Indent: Left: 0 cm, Hanging: 0,2 cm, Bulleted + Level: 1 + Aligned at: 0 cm + Indent at: 0,63 cm

Formatted

Formatted

Formatted

Formatted: Left

Formatted

Formatted: Font: 8 pt

Formatted: List Paragraph, Indent: Left: 0 cm, Hanging: 0,2 cm, Bulleted + Level: 1 + Aligned at: 0 cm + Indent at: 0,63 cm

Formatted

Formatted: List Paragraph, Indent: Left: 0 cm, Hanging: 0,2 cm, Bulleted + Level: 1 + Aligned at: 0 cm + Indent at: 0,63 cm

Formatted: Font: 8 pt

Formatted: Left

Formatted: Left

Formatted

Formatted

Formatted

Formatted: Left

Formatted: English (United States)

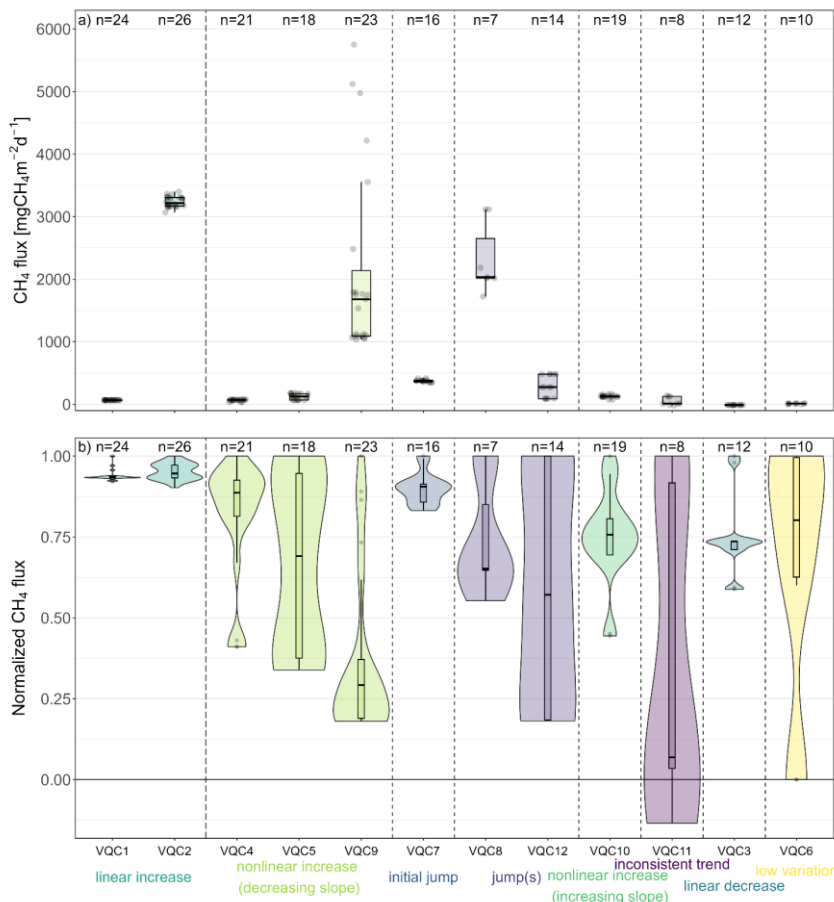


Table 1: VQC exercise (blue shading): Fluxes in mgCH₄m⁻²d⁻¹ calculated for each chamber measurement based on the time periods chosen for flux calculation by each survey participant. Coefficients of variance (CV) are given for each example measurement and measurement class across the participants. In the table, “NR” means that a participant did not respond to the respective measurement example so neither kept nor discarded the measurement. “D” indicates that a participant discarded a measurement example. Participants who gave no response to more than one measurement example are marked with “n.p.” for “not participated”.

Percentage of measurements kept for flux calculation (purple shading): Percentage of measurement examples within the visual QC exercise kept for flux calculation by participant (Kept_{QC} , relative to total number of measurement examples that a participant responded to), measurement example (Kept_{ex}), and measurement class ($\text{Kept}_{\text{class}}$). Extrapolation to Siikaneva data set and uncertainty estimates (red shading): Extrapolation of the visual QC results to the entire Siikaneva data set through weighting by the frequency of occurrence of each measurement class in the data set. Weighting factors are derived as the relative occurrence of the respective measurement class in the Siikaneva data set (788 measurement in total) divided by the number of occurrences in the visual QC exercise.

Measure- ment class	Linear increase		Nonlinear increase—decreasing slope			Initial jump	Jump(s)		Nonlinear increase — increase in slope	Inconsis- tent trend	Linear decrease	No trend
Weight [%]	468 (59.4)		144 (18.3)			66 (8.4)	62 (7.9)		25 (3.2)	16 (2.6)	4 (0.5)	3 (0.4)
Age-HD	VQC1	VQC2	VQC4	VQC5	VQC9	VQC7	VQC8	VQC12	VQC10	VQC11	VQC3	VQC6
Weight [%]	29.7	29.7	6.1	6.1	6.1	8.4	4.0	4.0	3.2	3.0	0.5	0.4

Run ID	Run Data												Kept acc [%]	Kept [%]
	VQC1	VQC2	VQC4	VQC5	VQC9	VQC7	VQC8	VQC12	VQC10	VQC11	VQC3	VQC6		
1	67.31	3204.96	68.56	171.47	1102.58	351.70	2183.91	87.58	137.75	127.25	-8.99	14.17	400	100
2	66.89	3169.76	Ø	Ø	1049.99	377.13	Ø	Ø	131.51	Ø	Ø	Ø	42	77
3	68.68	3367.11	73.95	169.61	1036.42	371.20	Ø	88.95	158.75	-18.64	-10.78	13.96	92	96
4	67.02	3294.94	66.92	171.47	1790.25	350.85	3117.04	482.53	127.40	Ø	-11.16	Ø	83	96
5	67.02*	3166.90*	32.80*	67.90*	1751.88*	Ø	Ø	Ø	116.73*	Ø	-11.16*	Ø	67	81
6	66.34	3156.77	65.24 [±]	61.26	1535.05	Ø	Ø	275.62	Ø	Ø	Ø	Ø	50	52
7	67.17	3067.51	61.86	Ø	1057.30	Ø	2031.93	Ø	133.38	Ø	Ø	Ø	50	70
8	67.02	3169.76	Ø	Ø	1091.37	378.93	Ø	Ø	116.73	Ø	-11.16	Ø	50	78
9	67.02	3169.76 [±]	71.13	67.90 [±]	1751.88*	Ø	Ø	Ø	Ø	Ø	-11.08	Ø	58	70
10	67.02	3304.26	75.34	Ø	Ø	380.34	Ø	Ø	158.75	Ø	Ø	Ø	42	77
11 (n-p)	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NA	NA
12 (n-p)	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NA	NA
13 (n-p)	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NA	NA
14	Ø	3166.90*	72.53	180.71	4216.13	Ø	Ø	Ø	168.07	Ø	Ø	Ø	42	51
15 (n-p)	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NA	NA
16	67.02*	3166.90*	72.72	110.49	3553.67	380.34	2016.94	482.53*	125.25	Ø	-11.16*	Ø	83	98
17 (n-p)	Ø	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NA	NA
18	66.14	3293.94	32.80*	67.90*	1122.55	357.19	Ø	482.53*	131.32	Ø	-9.58*	Ø	71	94
19	67.02	3169.76	34.38	69.94	1790.25	416.49	3117.04	482.53	116.73	9.58	-11.16	9.95	100	100
20 (n-p)	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NA	NA
21	67.31	3329.48	66.81	171.47	1084.14	357.19	Ø	88.36	125.19	127.25	-8.92	14.17	92	96
22	71.78	3201.49	79.90 [±]	67.90*	4973.26	Ø	Ø	Ø	75.82 [±]	Ø	Ø	Ø	50	81
23	66.34	3141.12	70.84	130.21 [±]	1680.02	371.20	2016.94	88.95	142.01	-6.17	-14.85	14.16	100	100
24	67.31	3299.19	73.94	169.61	5122.50	380.27	Ø	273.84	24.68	Ø	-10.76	Ø	75	94
25	Ø	Ø	Ø	Ø	Ø	Ø	Ø	Ø	NR	Ø	Ø	Ø	Ø	Ø
26	67.02	3204.96	76.38	Ø	Ø	Ø	Ø	Ø	Ø	Ø	-11.16	Ø	33	66
27 (n-p)	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NA	NA
28	67.02	3286.36	53.58 [±]	119.60	2482.08	Ø	Ø	275.62	Ø	Ø	Ø	Ø	50	82
29	67.17	3178.85 [±]	Ø	66.39 [±]	1764.55	377.13	1723.41	Ø	127.24	8.45	Ø	12.77	75	96

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Q0	D	3358.45	D	D	1088.47	346.32	D	D	D	D	D	8.51	33	45
Q1	67.02*	3312.29	72.72	180.71	5750.52	180.71*	D	482.53*	116.73*	138.74	D	D	75	95
Q2 (m.p.)	67.02*	3304.26	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	100	NA
Q3 (m.p.)	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NA	NA
Q4	69.67	3304.26	65.05	138.58	1065.04	380.34	D	275.62	D	D	-15.16	9.95	75	91
Q5 (m.p.)	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NR	NA	NA
Q6	67.31	3321.94	73.94	D	1124.44	D	D	88.95	130.74*	D	D	D	50	79
Mean±s	67.36±	3238.93±	65.07±	121.29±	2129.75±	374.37±	3315.31±	282.58±	126.57±	49.50±	-	9.76±	Weighted sum:	80±
D	1.18	84.93	4.52	8.72	90.04	9.88	64.43	72.84	3.50	8.31	11.36±	55		23
CV (%)	3	3	32	40	70	5	34	61	19	138	17	57		26
CV (%)	2.5		44			5	42.5		19	138	17	57		
CV (%)	86	96	81	69	88	62	27	54	76	29	50	38		
CV (%)	91		79			62	40		76	29	50	38		

* accepted but no time period was given for flux calculation. The flux was therefore estimated based on the entire measurement.
 † Flux was estimated based on a linear fit although the participant suggested to use a nonlinear model instead.

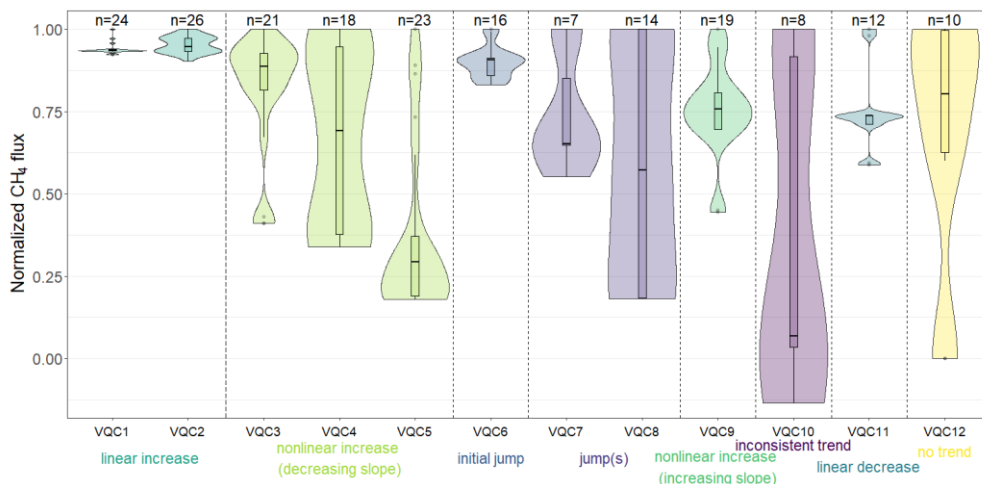


Figure 76: Distribution of researcher variability in flux estimates for each measurement example in the visual QC exercise (VQC1 – VQC12) by measurement scenario class. Range (a) and distribution (b) of flux estimates across the respondents. The number of survey participants (n) who contributed a flux estimate to the respective measurement example by selecting a time period for flux calculation is given on top of each boxplot (a) or violin (b), respectively. In (b), the flux estimates are normalized to the maximum flux estimate within each measurement example. The violins are scaled to all have the same maximum width. Violins crossing the zero line indicate that, for the respective measurement example, the selection of the time period for flux calculation made the difference between CH₄ emission and uptake.

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

~~The number of survey participants (n) who contributed a flux estimate to the respective measurement example by selecting a time period for flux calculation is given on top of each violin.~~

3.2.2.2 Linear increase

Two example measurements included in the visual QC exercise showed a linear increase in CH₄ concentrations over the entire time of the chamber closure. In the first example (measurement ID VQC1, Figure 7), the CH₄ concentrations increased by only 0.4 ppm from a starting concentration of 2.1 ppm to a concentration of 2.5 ppm at the end of the measurement while in the second example (measurement ID VQC2, Figure 8) the starting concentration of 3.8 ppm was higher and increased by 38.2 ppm to reach 42.0 ppm over the course of the measurement. The CO₂ concentration in the chamber decreased during both measurements by 13.4 ppm and 9.1 ppm, respectively. The starting concentration of H₂O_A in VQC2 was more than 10 times higher than for VQC1 and decreased strongly over the course of the measurement with an abrupt decrease at around 50 s after chamber closure, while H₂O_A concentrations increased slightly in VQC1.

The participants described the trend in CH₄ concentrations in both VQC1 and VQC2 as a linear increase which they explained by net CH₄ production and diffusive emission. The CH₄ flux in VQC2 was additionally classified as large with one participant concluding that the measurement plot was a “hotspot” for CH₄ emission. For both VQC1 and VQC2, some participants additionally noticed slight deviations from the linear behaviour of the CH₄ concentrations. Minor jumps in the CH₄ concentration in VQC1 were mentioned by 17 participants (61%), which they related to CH₄ ebullition (9 participants), insufficient mixing due a defective fan (3 participants), wind (1 participant), wind induced pressure changes (1 participant), changes in atmospheric pressure influencing the ground diffusion rates and/or atmospheric pressure gradient (1 participant), boundary layer disturbance (1 participant), leakage (2 participants), disturbance (1 participant) caused by chamber placement or footsteps (1 participant). For VQC2, half of the participants pointed out a decrease in the slope of CH₄ concentrations starting between 250 and 260 s after the chamber closure, 21% of whom also noticed a simultaneous decrease in the slope of CO₂ concentrations. As explanations the participants mentioned saturation of the chamber headspace decreasing the concentration gradient over time (5 participants), a build up of pressure (2 participants) potentially due to a defective pressure valve towards the end of the measurement (1 participant), a change in temperature over the course of the measurement (1 participant), or a small leak (1 participant) probably combined with windy conditions (1 participant). Many participants furthermore discussed the change in CO₂ and H₂O_A concentrations over the time of the chamber closure. For VQC1, three participants mentioned that the CO₂ and H₂O_A concentrations show a linear change, two of whom concluded that there was no air leaking from the chamber. Three participants on the other hand were concerned about the H₂O_A measurements due to the high and increasing concentrations, and due to an assumed saturation and therefore decreasing slope towards the end of the measurement. Leakage from the chamber was suspected by three participants, two of whom explained this presumption with vegetation overgrowing the collar and one with the use of a less airtight rubber seal as opposed to a water seal. For VQC2, 18% of the participants picked up on the drop in H₂O_A concentrations occurring around 40 s after the chamber closure, 40% of whom additionally mentioned a simultaneous change in the slope of CO₂ concentrations. Their reasoning included water condensing on the chamber walls and changing light conditions. Few participants decided to discard the two measurement

Formatted: English (United States)

Formatted: Keep with next, Keep lines together

Formatted: English (United States)

Formatted: English (United States)

Formatted: Indent: First line: 0 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

820 examples. Measurement VQC1 was discarded by 4 participants (14%) suspecting CH₄ ebullition or stating that the starting
concentrations of CO₂ were too high above ambient concentrations or that all chamber measurements generally need to be
shaded. One participant excluded VQC2 due to an assumed saturation effect and one additional participant mentioned
825 ebullition and a high initial concentration of CH₄ as potential reasons to exclude the measurement from flux calculations. 86%
and 89% of the participants decided to keep VQC1 and VQC2 for flux calculation, respectively, due to the consistent linear
increase in CH₄ concentrations without clear indications of significant disturbances or any malfunctioning of the instruments.
For VQC1, the participants further supported their decision with the linear change in CO₂ and H₂O₂ concentrations making
leakage from the chamber unlikely as well as with near ambient CH₄ concentrations at the measurement start. For both VQC1
and VQC2 most participants who gave start and end times for flux calculation chose the middle part of the measurement,
discarding the beginning and the end without mentioning a specific reason. The remaining participants considered the CO₂
830 and/or H₂O₂ concentrations in their choice of the time period for curve fitting. For VQC1, three participants chose the beginning
of the measurement only, resulting in slightly higher flux estimates, two of whom assumed that H₂O₂ saturation diminished the
increase in CH₄ concentrations towards the end of the measurement. For VQC2, some participants acknowledged the strong
drop in H₂O₂ concentrations. Having no further information on potential reasons three of them decided not to let this unexpected
behaviour in H₂O₂ concentrations make them discard the CH₄ measurements while other participants reacted by excluding the
835 time of the drop in H₂O₂ concentrations from their calculation of the CH₄ flux through either using the part of the measurement
after the drop (7 participants) or before the drop (1 participant). 61% of the 23 participants who entered start and end times for
flux calculation discarded the end of the measurement where CH₄ and CO₂ concentrations increased at a lower rate, resulting
in slightly higher flux estimates above 3200 mgCH₄m⁻²d⁻¹. Two participants suggested to use a nonlinear fit which one of
them specified as exponential.

840

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

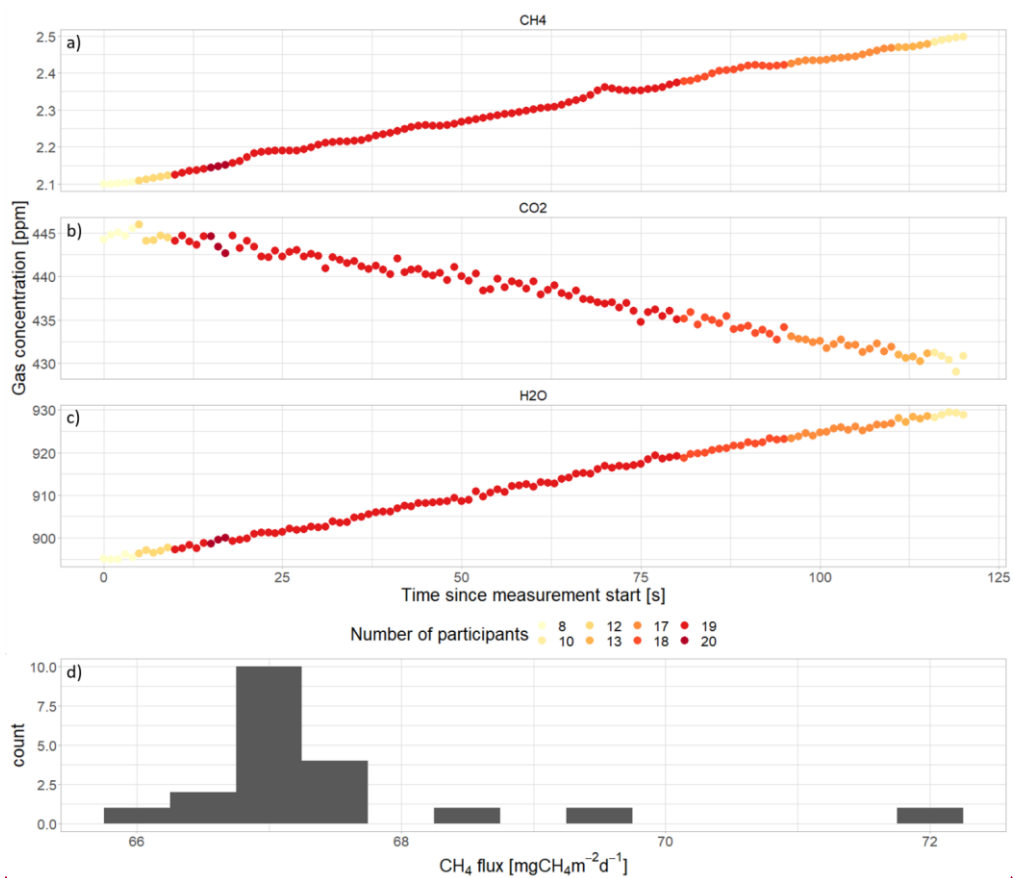


Figure 7: Measurement example VQC1 of a small linear increase in CH₄ concentrations over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colours of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

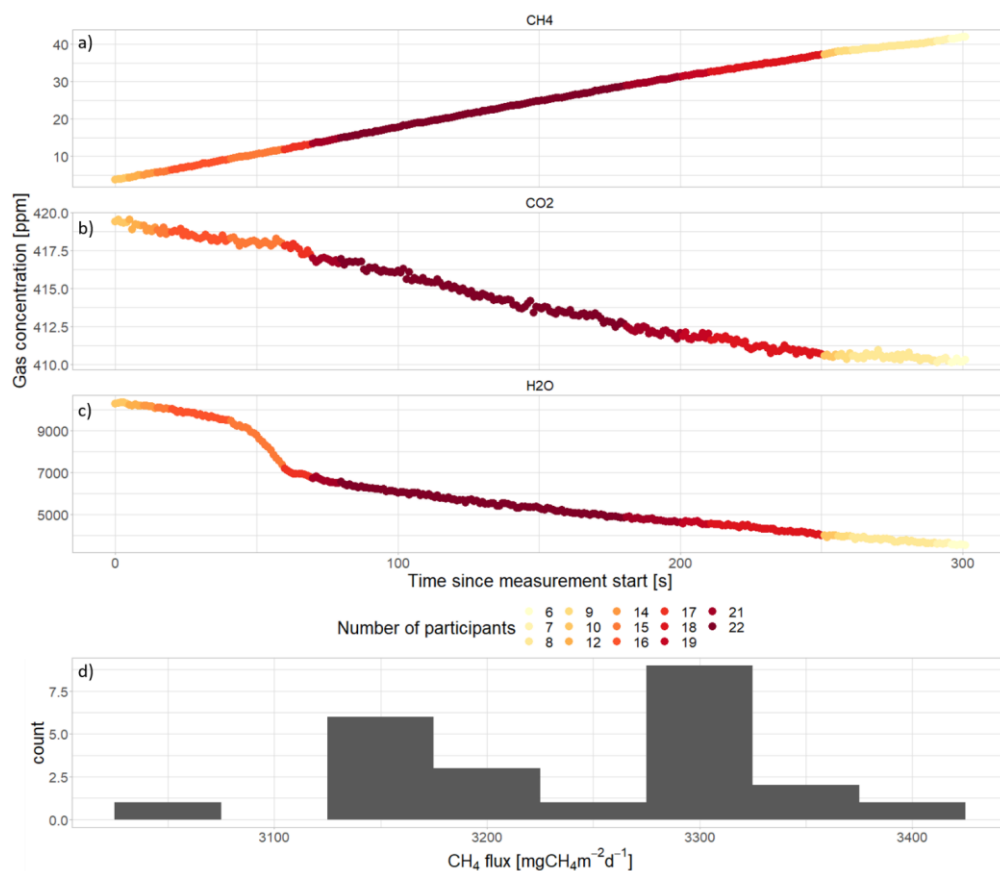


Figure 8: Measurement example VQC2 of a strong linear increase in CH₄ concentrations over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colours of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

3.2.3 Nonlinear increase—decreasing slope

In the visual QC exercise, we included three examples of measurements that feature a nonlinear increase in CH₄ concentrations during the chamber closure with the rate of increase flattening out over time. Two examples show a small nonlinear increase in CH₄ concentrations (measurement IDs VQC4 and VQC5, Figures 9 and 10) simultaneous with linearly decreasing CO₂

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: Normal

concentrations. H_2O concentrations increased over the time of the chamber closure in VQC4 but decreased in VQC5. The third example (measurement ID VQC9, Figure 11) shows a stronger increase in CH_4 concentrations with intermittent jumps, linearly increasing CO_2 concentrations and H_2O concentrations that fluctuate without a clear trend.

The participants classified the CH₄ measurements in VQC5 as a small flux that resulted from a balance between CH₄ production and oxidation while VQC9 was identified as large emission indicating a CH₄ hotspot. The majority of the participants (85%, 85%, and 81%) discussed the nonlinear behaviour of the CH₄ concentrations in VQC4, VQC5, and VQC9, respectively, offering various explanations for the decreasing rate of increase over time that were mainly related to chamber saturation, chamber leakage or an initial disturbance (Table 2).

Most participants (9, 10, and 8) suspected a saturation of the chamber headspace, while two participants stated that saturation was unlikely to be reached during a measurement as short as VQC4 (330 seconds) and one participant explicitly mentioned that the changing slope in VQC9 did not look like a saturation effect. Just as many participants (9) suggested leakage through a weakening seal as the reason for the decreasing slope in VQC4, as supported by the simultaneously decreasing slope in CO_2 and H_2O_2 concentrations while other participants explicitly stated that CO_2 and H_2O_2 concentrations did not indicate a leak in this measurement. Due to the consistently linear CO_2 concentrations in VQC5 and VQC9, only one participant each suspected leakage during these measurement examples. For VQC4, three participants further suspected that the high H_2O_2 concentrations at the end of the measurement influenced the CH_4 measurements, for example through condensation inside the chamber or in the gas flow line, one participant suggested a varying performance of the chamber fan, and two participants assumed that the nonlinearity was a phenomenon specific to *Sphagnum* moss. One participant more generally suggested that vegetation effects changed over the course of the measurement due to changing light conditions, affecting the CO_2 and H_2O_2 concentrations in VQC4 and the CH_4 concentrations in VQC5.

Besides a saturation effect or a weakening seal that would cause a decreasing slope in CH_4 concentrations towards the end of the measurement, many participants (3, 6 and 8) suggested that an initial disturbance such as ebullition triggered by the chamber placement had caused the stronger increase in the beginning of measurement examples VQC4, VQC5, and VQC9. For VQC9, 31% of the participants additionally pointed out minor fluctuations superimposed on the overall nonlinear increase in CH_4 concentrations. Two thirds of them referred to the fluctuations as minor ebullition events while the others suggested episodic leakage from the chamber potentially caused by gusts of wind lifting the chamber sides or a malfunctioning pressure gauge. One participant pointed out that the CH_4 fluctuations cooccurred with fluctuations in the H_2O concentrations and therefore suspected an instrument issue that could be related to spikes in the instrument cavity pressure.

The nonlinearity in the CH₄ concentrations resulted in 15%, 31%, and 12% of the participants deciding to discard the entire measurement example VQC4, VQC5, and VQC9, respectively (Table 2). The reasons mentioned for the exclusion of the measurements again reflected the different interpretations of the participants on which part of the measurement represented the real flux. This disagreement shows less strongly in the range of flux estimates since participants who suspected an initial disturbance of the measurement disproportionately often discarded the entire measurement as they assumed that an initial disturbance would also affect the remaining part of the measurement. For VQC4 and VQC5, all 54% of the participants who

Formatted: English (United States)

Formatted: English (United States)

Formatted: Indent: First line: 0 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

895

900

905

provided start and end times for flux calculations agreed that the beginning of the measurement should be used for or at least be included in the flux calculation, with three participants suggesting a nonlinear fit for both measurement examples. This resulted in smaller ranges of flux estimates compared to VQC9 (Table 2) which instead reflects the fundamentally different interpretations among the participants on which part of the measurement should be used for flux calculation. Here, half of the 21% of participants who gave start and end times for flux calculation chose a later part of the measurement where CH₄ concentrations appeared linear over a longer time period. This resulted in lower flux estimates (between 1000 and 1200 mgCH₄m⁻²d⁻¹) compared to the flux estimates larger than 3500 mgCH₄m⁻²d⁻¹ derived for the one quarter of participants who instead chose the beginning of the measurement (Figure 11).

Table 2: Explanations for the nonlinear increase in CH₄ concentrations, reasons to discard, and reasons and ways to keep measurements showing an increase in CH₄ concentrations as given by the participants. The responses were categorized based on the free text entries for measurement examples VQC4, VQC5, and VQC9. The number of responses given in the respective category are provided in brackets.

Explanations for nonlinearity	Reasons to discard	Reasons to keep
Saturation (23)	Saturation (2) Nonlinearity—no steady state reached (3)	A nonlinear fit can be used (9)
Initial disturbance (16)	Initial disturbance biases flux later on (2)	A (linear) part of the curve can still be used (41)
Bad seal / Leakage from the chamber (8)	Bad seal / Leakage from the chamber (4)	
Unsure (12)	Unclear which part of the measurement represents the real flux (3)	No clear disturbance of the measurement (9)
Changing environmental conditions (1)	Changing environmental conditions (1)	Linear trend in CO ₂ concentrations (5)

Formatted: English (United States)

Formatted: English (United States)

Formatted: Justified

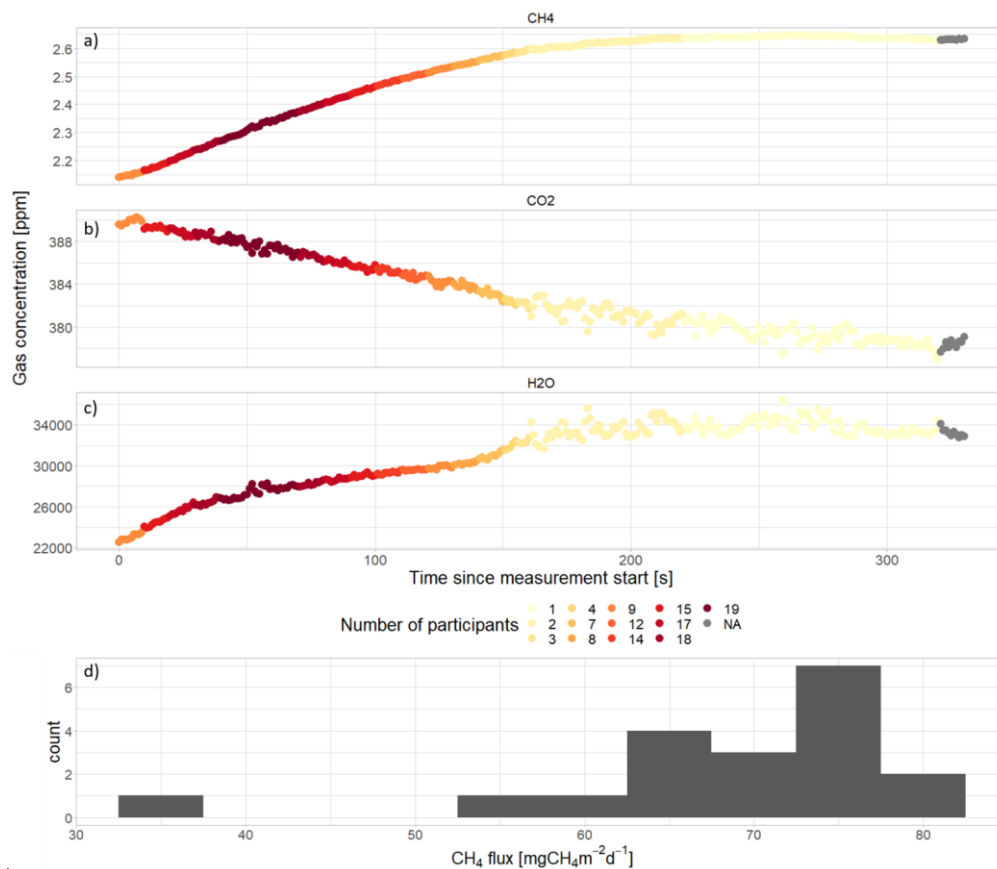


Figure 9: Measurement example VQC4 of a small nonlinear increase in CH₄ concentrations with decreasing slope over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colours of the data points indicating how many participants included the respective data point in the time period that they chose for flux

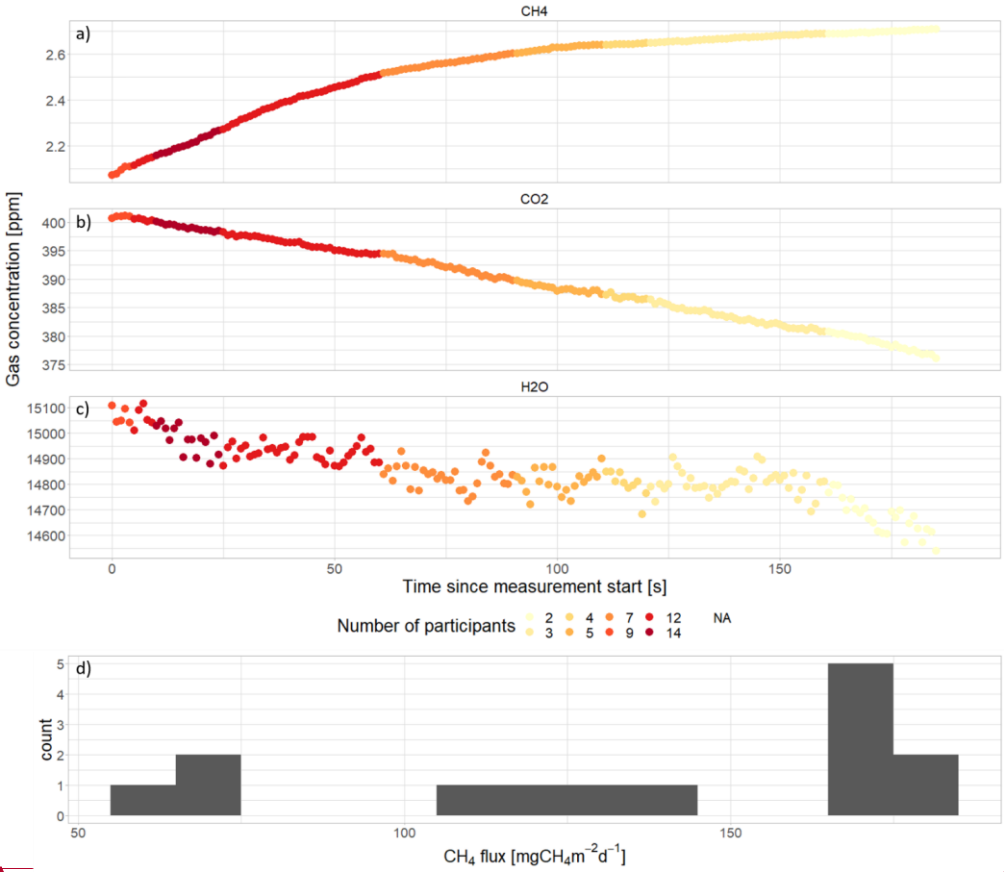
Formatted: Font: Not Italic, English (United States)

Formatted: Heading 1

Formatted: Font: Not Bold, English (United States)

Formatted: Font: Not Bold, English (United States)

calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).



915 **Figure 10:** Measurement example VQC5 of a small nonlinear increase in CH₄ concentrations with decreasing slope over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colours of the data points indicating how many participants included the respective data point in the time period that they chose for flux

Formatted: English (United States)
Formatted: Font: Not Italic, English (United States)

Formatted: Font: Not Bold, English (United States)
Formatted: Font: Not Bold, English (United States)

calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

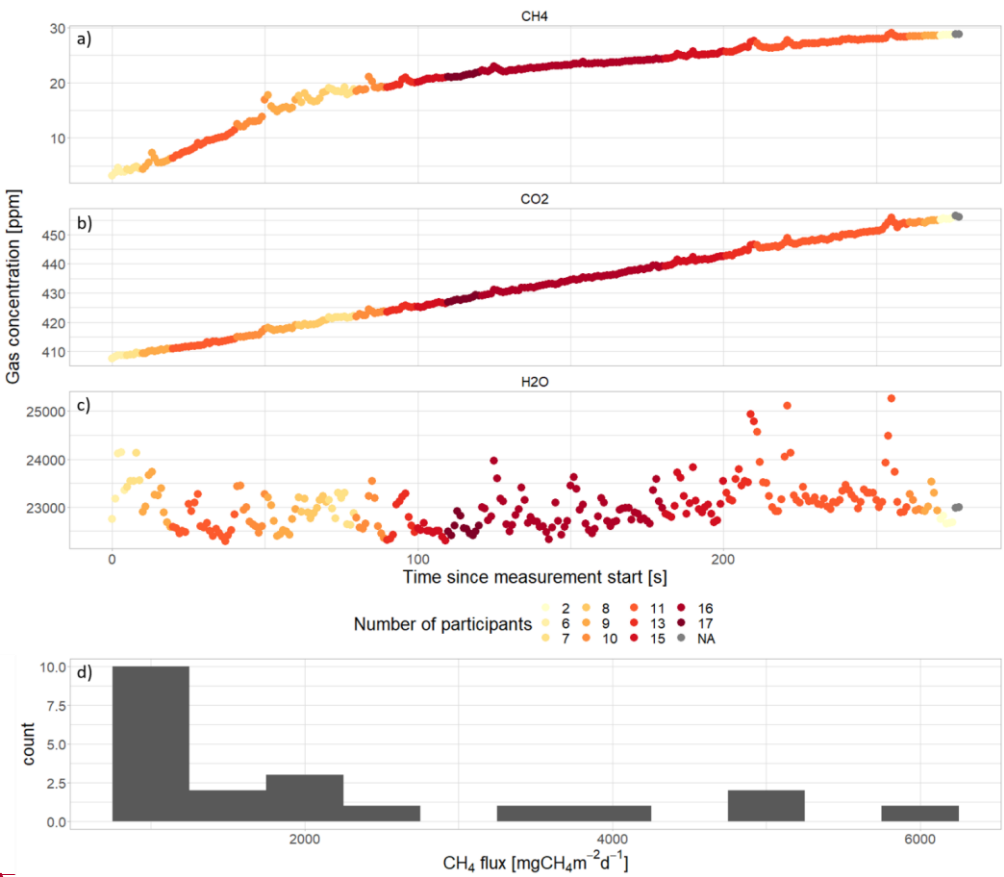


Figure 11: Measurement example VQC9 of a strong nonlinear and jumpy increase in CH₄ concentrations with decreasing slope over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colours of the data points indicating how many participants included the respective data point in the time period that they chose for flux

Formatted: Font: Not Italic, English (United States)

Formatted: Font: Not Bold, English (United States)

Formatted: Font: Not Bold, English (United States)

925 calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram
of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

930 **3.2.4 Jump(s) at the beginning or in the middle of measurements**

In our visual QC exercise, we included three example measurements that showed a relatively linear increase in CH₄ concentrations that was interrupted by one (VQC8, Figure 12 and VQC7, Figure 13) or more (VQC12, Figure 14) sudden increases in the concentration. In one, this occurred at the beginning (VQC7) and in the others, the middle of the measurement (VQC8, VQC12). In examples VQC8 and VQC7, these sudden jumps appeared in all three gases with CH₄ and H₂O concentrations showing a sudden increase while CO₂ concentrations dropped simultaneously. In VQC12, on the contrary, CO₂ and H₂O showed no equivalent to the jumps in the CH₄ concentration. In VQC8, a strong decrease in CH₄ concentrations directly followed the sudden increase, while in VQC7 and VQC12 the concentrations continued to increase at a lower rate starting close to the high concentration level after the jump.

Nearly all (100%, 65%, and 92%) of the participants mentioned the jump(s) in CH₄ concentration when discussing the measurement examples VQC8, VQC7, and VQC12, respectively. For all measurement examples, the majority of these participants explained their observation with episodic events of ebullitive CH₄ emission (VQC8: 65%, VQC7: 88%, VQC12: 92%) with only one participant each suggesting a malfunctioning of the gas analyser as a reason for the sudden increase in CH₄ concentrations in VQC7 and VQC12. Some (35%) of the participants assuming ebullition stated that the ebullition event in VQC8 was caused by a disturbance and all agreed that the chamber placement caused the ebullition for VQC7. Only one participant (5%) mentioned anthropogenic disturbance as the reason for the ebullition events in VQC12. For VQC8, 12% of the participants pointed out the sudden changes in CO₂ and H₂O concentrations along with the jumps in CH₄. Reasons mentioned by one participant each were a malfunctioning of the gas analyser and an overpressure caused by the bubble release while another participant suggested the release of gas bubbles with high CH₄ but low CO₂ concentrations as a natural cause for this observation. Similarly, one of the two participants who mentioned the absence of a simultaneous change in the other gases in VQC12, assumed a release of bubbles with high CH₄ concentration but CO₂ concentrations close to ambient conditions due to the different production depths of the two gases. For VQC8, 41% of the participants discussed the decrease in CH₄ concentrations following the assumed ebullition event and suggested leakage of air from the chamber, potentially combined with wind as a potential cause. In the discussion of VQC7, two participants disagreed on the effect of the water table on CH₄ ebullition, one mentioning that in the measurement CH₄ ebullition was more likely to happen because of the high water table while the other stated that ebullition happened despite the high water table, indicating a fundamentally different understanding of the causes of CH₄ ebullition among the participants. Two participants of VQC7 furthermore classified the measurement as an example of strong CH₄ emission which they explained by strong anaerobic CH₄ production related to the high water table and by the vegetation providing substrate for acetoclastic CH₄ production, respectively.

Of the three measurements with jumps in CH₄ concentrations that we included in the visual QC exercise, VQC8 raised the most concern with the highest number of participants excluding the example (Table 2) and with the largest variety of reasons mentioned for the discard, including the inconsistent trend in CH₄ and CO₂ concentrations making them wonder which part of the measurement to use for flux calculation, ebullition affecting the pressure inside the chamber, too much variation in CH₄ and CO₂ concentrations even after the jump, chamber leakage and too high initial CH₄ and CO₂ concentrations. Leakage was also suggested by one participant for VQC7, who suspected that Sphagnum moss

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: Heading 1, Indent: First line: 0 cm

Formatted: English (United States)

might have obstructed the chamber seal with the collar. VQC12 was classified as too short of a measurement by one participant and discarded by another for too high initial CO₂ concentrations.

There was disagreement among the participants on whether the remaining part of a measurement after a jump in the CH₄ concentration could still be used for flux calculation. For 10 of 11 participants discarding VQC7, the main concern was high concentrations having a lasting effect on the concentration gradient and thus on the diffusive CH₄ flux during the rest of the measurement while only one of 21 and 15 participants discarded measurements VQC8 and VQC12 for that reason. For VQC12, four of the 11 participants who kept the measurement, all of whom also gave start and end times, avoided this problem by using the beginning of the measurement before the first jump for flux calculation. On the contrary, for VQC8 and VQC7 five and 14 of the seven and 15 participants who kept the measurement and/or gave start and end times for flux calculation decided that the measurement after the jump in CH₄ concentrations could still be used for flux calculation, respectively, and five participants in VQC12 preferred to use the part between the first two jumps because it showed a longer linear increase.

The choice of different time periods for flux calculation resulted in two and three different classes of flux magnitudes for VQC8 and for VQC7 and VQC12, respectively. The highest flux estimates of more than 3000 mgCH₄ m⁻² d⁻¹, 483 mgCH₄ m⁻² d⁻¹, and 416 mgCH₄ m⁻² d⁻¹ stemmed from the two, one, and two participants who used the whole measurement example VQC7, VQC8, and VQC12, respectively, for flux calculation because these estimates also included ebullitive in addition to diffusive CH₄ emissions, reflecting the general disagreement on whether CH₄ ebullition should directly be included in the flux estimates derived from chamber measurements. For VQC7 and VQC12, the flux estimates from the participants who excluded the jumps in CH₄ concentration from the time period for flux calculation can further be split into two classes. For VQC7, nine participants excluded only the very beginning of the measurement, while five participants only used a later part starting at about 50 s into the measurement when CO₂ concentrations decreased at a higher rate, resulting in slightly lower CH₄ fluxes. For VQC12, when excluding the jump in CH₄ concentrations the flux estimates were higher for four participants who chose the measurement period before the first jump, reaching up to 275 mgCH₄ m⁻² d⁻¹ compared to the five participants who chose the longest linear part of the measurement leading to flux as low as 88 mgCH₄ m⁻² d⁻¹. Due to the very linear behaviour of the CH₄ concentrations

following the initial jump and the higher agreement on the time period used for flux calculation, the CV of 5% for VQC7 was much lower than for the CVs of 24 and 61% for VQC8 and VQC12, respectively.



Figure 12: Measurement example VQC8 of an overall increase in CH₄ concentrations over the time of the chamber closure after a strong jump in CH₄ concentrations. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colours of the data points indicating how many participants included the respective data point in the time period that they chose for

Formatted: Font: Not Italic, English (United States)

Formatted: Heading 1

Formatted: Font: Not Bold, English (United States)

Formatted: Font: Not Bold, English (United States)

995 flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

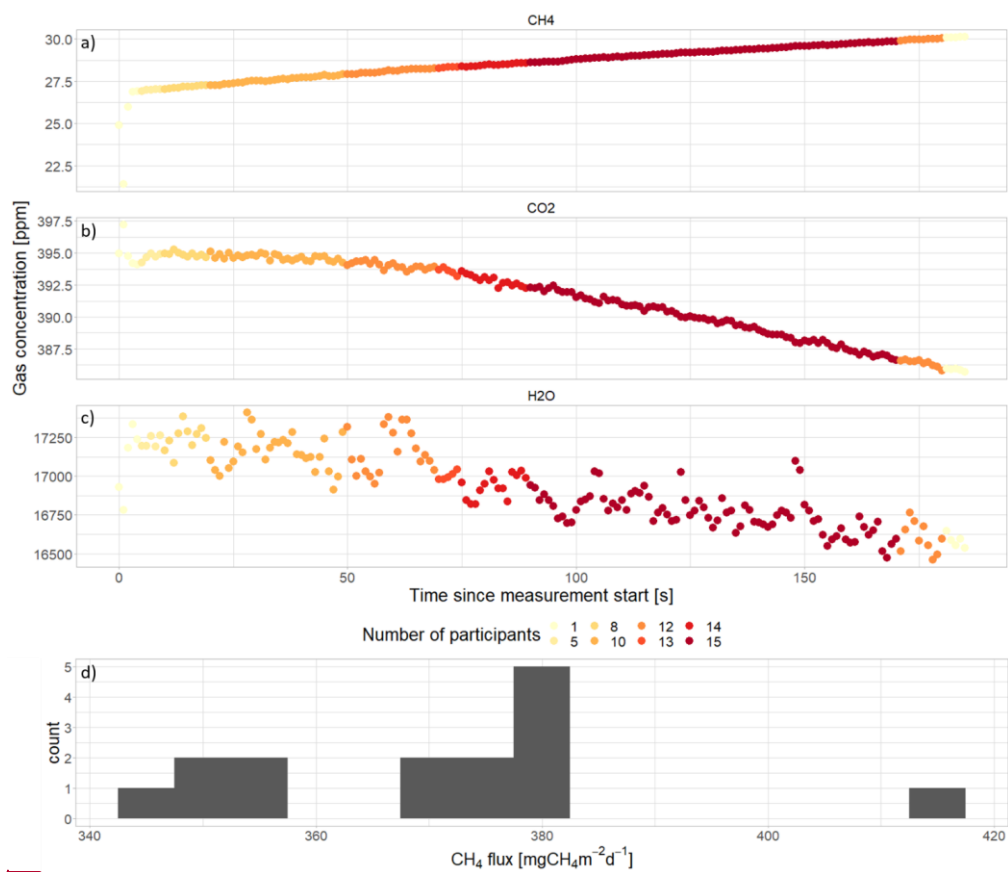


Figure 13: Measurement example VQC7 of a linear increase in CH₄ concentrations of the chamber closure after an initial jump in CH₄ concentrations. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colours of the data points indicating how many participants included the respective data point in the time period that they chose for flux

Formatted: Font: Not Italic, English (United States)

Formatted: Font: Not Bold, English (United States)

Formatted: Font: Not Bold, English (United States)

calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time-period chosen for flux calculation by the participants (d).

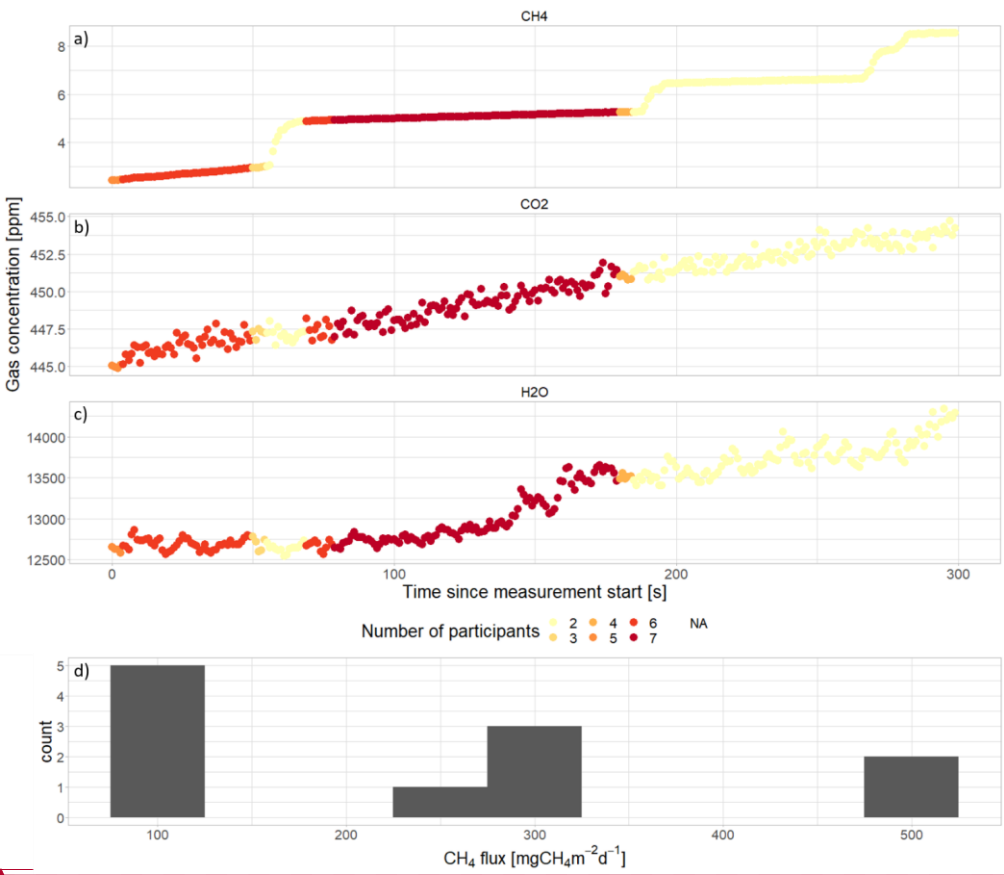


Figure 14: Measurement example VQC12 of a linear increase in CH₄ concentrations between repeated jumps in the CH₄ concentration over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colours of the data points indicating how many participants included the respective data point in the time period that they

Formatted: Font: Not Italic, English (United States)

Formatted: Font: Not Bold, English (United States)

Formatted: Font: Not Bold, English (United States)

chosed for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

3.2.5 Nonlinear increase ~ increasing slope

One example included in the visual QC exercise showed a nonlinear increase in CH₄ concentrations over the chamber closure with the rate of increase becoming stronger over time (measurement ID VQC10, Figure 15). 15% of the participants classified the measurement as a diffusive emission of CH₄ without mentioning further details while 65% discussed the increasing slope in CH₄ concentrations over time, suggesting various reasons that could have caused the observed shape of the curve. The reasons suggested included an initial period of mixing or adjusting, an increase in chamber temperature over time, a disturbance of the measurement plot, a disturbance of the concentration gradient in the soil during chamber placement, an influence of the chamber on plant-mediated CH₄ transport, an incomplete seal of the chamber, incomplete mixing, and an interference with the simultaneously increasing H₂O concentrations. Two participants mentioned that they had not seen such a shape in CH₄ concentrations from chamber measurements before. Regarding the magnitude of CH₄ emissions, three participants pointed out the strong increase in CH₄ concentrations despite the relatively low water table, which they related to plant-mediated CH₄ transport. One participant further mentioned that also the emission of CO₂ was high, indicating warm peat conditions. Two participants mentioned the higher and decreasing CO₂ concentrations in the beginning of the measurement which one of them related to the chamber placement, pushing more gases out of the ground. One participant furthermore mentioned that the chamber seal seemed to be intact.

Six participants decided to discard the measurement, three of whom did so because they could not explain the shape of the curve and stated that the curvature was so strong that the flux estimate would strongly depend on the time period chosen for flux calculation. The three remaining participants mentioned similarly unexpected shapes of CO₂ and H₂O concentrations, higher H₂O concentrations towards the end of the chamber closure which might have interfered with the CH₄ measurements, and high initial CH₄ concentrations as reasons to discard the measurement. 19 participants kept the measurement for flux calculation. The flux estimates for the 17 participants who gave start and end times for flux calculation strongly depended on the time period they chose which in turn depended on their interpretation of the measurement resulting in three distinct classes of flux magnitudes. Two participants decided to use the entire measurement, resulting in intermediate flux estimates of 117 mgCH₄.m⁻².d⁻¹. The majority (13) decided to remove the first 20 to 120 s of the measurement to keep only the more linear part of the CH₄ concentrations in the end, resulting in the highest flux estimates between 125 and 170 mgCH₄.m⁻².d⁻¹. The two remaining participants chose only the linear first 60 or 70 s of the measurement for flux calculation resulting in lower flux estimates of 75 and 76 mgCH₄.m⁻².d⁻¹.

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: Heading 1, Indent: First line: 0 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

respectively, due to the lower rate of increase. Two participants suggested to use a nonlinear fit which one of them specified as exponential.

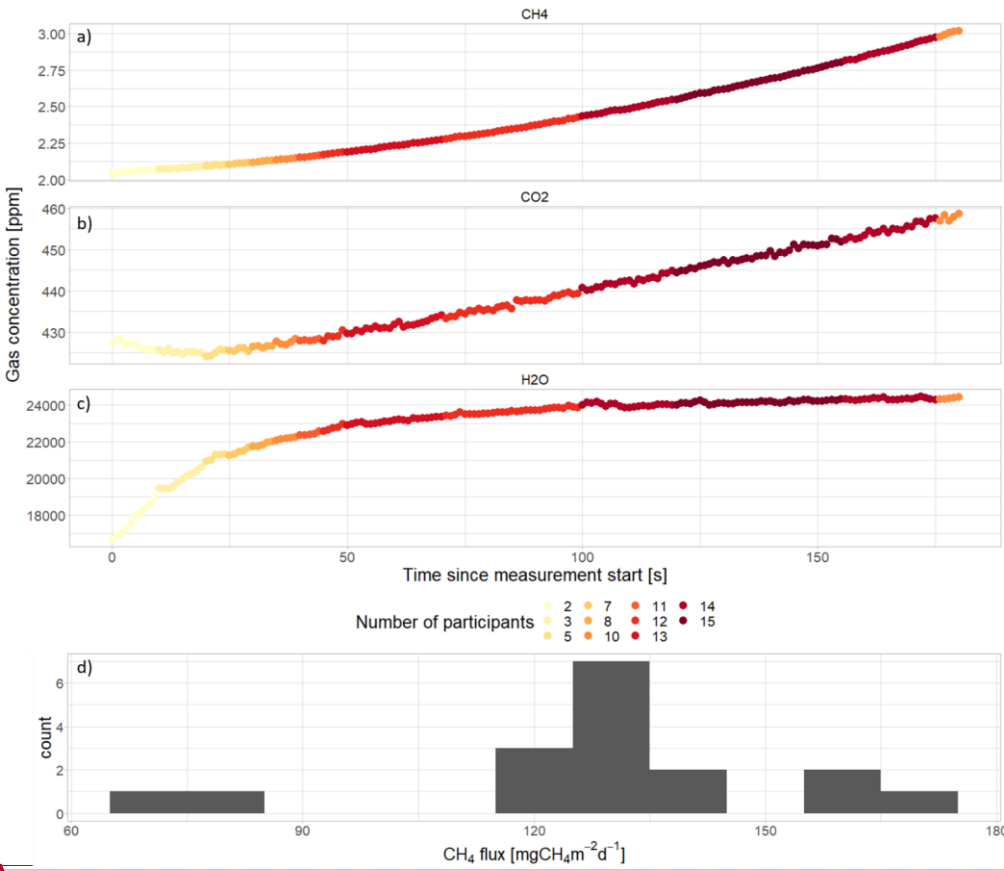


Figure 15: Measurement example VQC10 of a small nonlinear increase in CH₄ concentrations with increasing slope over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colours of the data points indicating how many participants included the respective data point in the time period that they chose for flux

Formatted: Font: Not Italic, English (United States)

Formatted: Heading 1

Formatted: Font: Not Bold, English (United States)

Formatted: Font: Not Bold, English (United States)

Formatted: Font: Not Bold, English (United States)

1045 calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram
of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

3.2.6 Inconsistent trend

1050 One example included in our visual QC exercise showed an inconsistent trend in CH₄ with a change from increasing to
decreasing concentrations over the time of the chamber closure (measurement ID VQC11, Figure 16). The survey
participants disagreed on the reason for this behaviour of the CH₄ concentrations. One part of the participants stated
that CH₄ oxidation as indicated by the decrease in CH₄ concentrations towards the end of the measurement was
unexpected and suggested that measurement issues were responsible for the inconsistent trend in CH₄ concentrations.
1055 They had different opinions however on the timing of the disturbance and therefore on which part of the measurement
represented the actual CH₄ flux. Some participants suggested an initial disturbance such as CH₄ ebullition caused by
the chamber placement while others assumed that the measurement was disturbed at a later point by a problem with
the CH₄ analyzer like saturation of the detector or H₂O interference due to the high concentrations towards the end of
the measurement and potentially condensation of water vapor, or leakage or a malfunctioning fan after about 50 s into
the measurement.

1060 Most participants (66%) discarded the measurement because they missed a consistent trend of sufficient length in the
CH₄ concentrations. Since the changing trend was either related to a disturbance or the reason was described as
unclear, the participants did not know which part of the measurement to use for the flux calculation. Two participants
additionally discarded the measurement because they considered the changes in the CH₄ concentration as too close to
zero and another participant mentioned that the CO₂ and H₂O concentrations did not show a steady trend over time
1065 either. Some (23%) of the participants decided to keep the measurement for flux calculation, all of whom provided
start and end times for flux calculation as well as one additional participant who was uncertain whether to keep or to
discard the measurement. The choice of the time periods used for flux calculation depended on the interpretation of
the observed pattern in CH₄ concentrations and thus strongly influenced the resulting flux estimate ranging between a
CH₄ uptake of -19 mgCH₄ m⁻² d⁻¹ to CH₄ emissions of up to 139 mgCH₄ m⁻² d⁻¹ and splitting the flux histogram into
1070 three distinct modes. Two participants chose to keep the entire measurement, resulting in a small positive flux
indicating small net CH₄ emission of 8 to 10 mgCH₄ m⁻² d⁻¹. Three participants decided to use the stronger increase in
CH₄ concentrations in the beginning of the measurement, resulting in the highest CH₄ emissions between 127 and
139 mgCH₄ m⁻² d⁻¹ while two participants assumed that CH₄ was consumed at the plot, using the later decreasing part

Formatted: English (United States)

Formatted: English (United States)

Formatted: Heading 1, Indent: First line: 0 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

075

of the CH_4 concentrations, resulting in negative flux estimates between -6 and $-19 \text{ mgCH}_4 \text{ m}^{-2} \text{ d}^{-1}$. This resulted in the highest CV among the measurement classes, estimated at 138%.

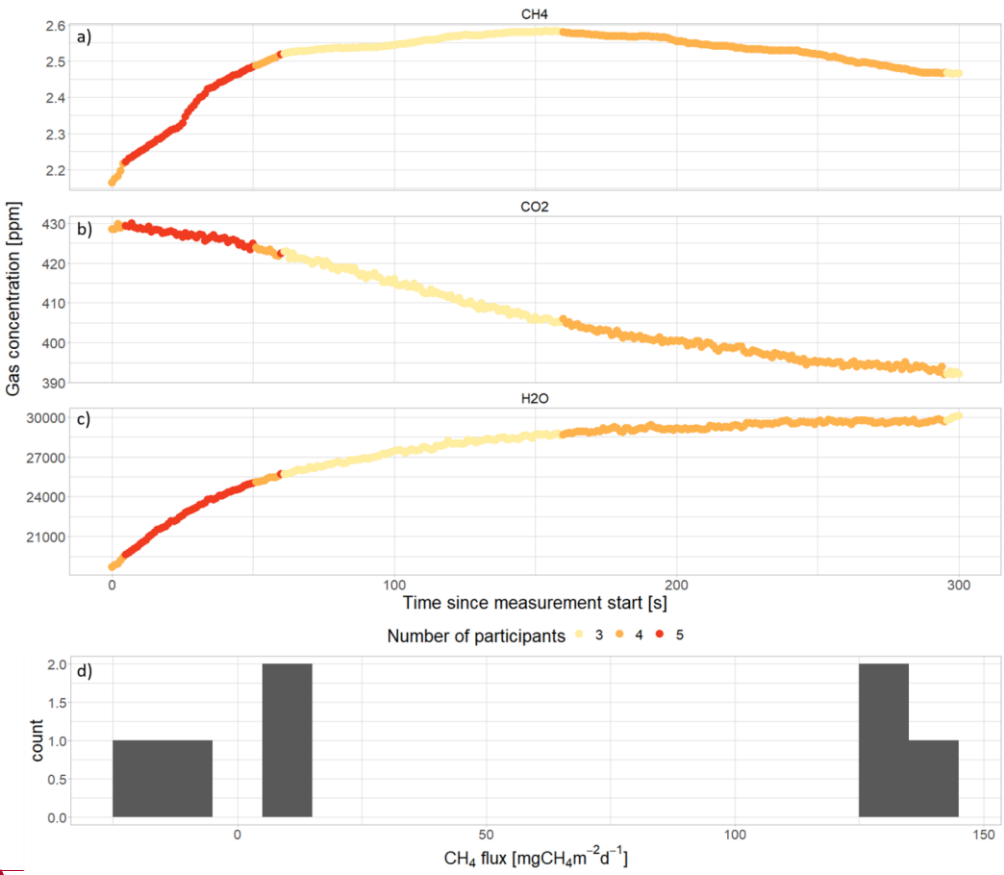


Figure 16: Measurement example VQCH of CH_4 concentrations showing an inconsistent trend over the time of the chamber closure. Simultaneous measurements of CH_4 (a), CO_2 (b), and H_2O (c) concentrations over time during chamber closure with the colours of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the

Formatted: English (United States)

Formatted: English (United States)

Formatted: Font: Not Italic, English (United States)

Formatted: Heading 1

Formatted: Font: Not Bold, English (United States)

Formatted: Font: Not Bold, English (United States)

1080 responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes
calculated based on the time period chosen for flux calculation by the participants (d).

3.2.7 Linear decrease

1085 One of the measurements in the visual QC exercise showed a small linear decrease in CH₄ concentrations over time
(measurement ID VQC3, Figure 17). The survey participants largely disagreed on whether this measurement
represented a real CH₄ flux. The majority (65%) of the participants assumed real net CH₄ uptake due to CH₄ oxidation
dominating over CH₄ production while some (19%) of the participants referred to leakage and too high initial CH₄
1090 concentrations in the chamber as technical problems causing a false apparent uptake of CH₄. The remaining 15% of
the participants explicitly stated that they were unsure if the measurement represented a real flux. 23% of the
participants more specifically mentioned an inconsistent trend in the CH₄ concentrations referring to three different
stages of CH₄ flux or nonlinearities at the beginning and at the end of the measurement. As explanations, they offered
initial CH₄ ebullition caused by the chamber placement, changes in the chamber temperature, changes in wind speed
combined with chamber leakage, or changes in PAR potentially due to a changing cloud cover or due to condensation
1095 inside the chamber indicated by the trend in CO₂ concentrations changing along with the CH₄ trend as well as by high
H₂O concentrations.

A slim majority (54%) of the participants discarded the measurement because they did not expect CH₄ uptake in the
given environmental (despite the relatively low water table), or because of the inconsistent trend in CH₄ concentrations
which makes them unsure which part of the measurement to use for flux calculation, or because of too high initial
concentrations of CH₄ and/or CO₂, or because they suspected anthropogenic disturbance from footprints and
1100 compacted vegetation or leakage. The flux estimates derived from the start and end times given by 11 of the 12
participants who decided to keep the measurement (46%) differed between the time periods chosen for flux calculation.
While five participants chose the entire measurement, resulting in intermediate values of CH₄ uptake, the remaining
six participants chose the time period for curve fitting based on the CO₂ concentrations. The middle part of the
measurement with linearly decreasing CO₂ concentrations, the beginning of the measurement with stable CO₂
1105 concentrations, and the end of the measurement with linearly increasing CO₂ concentrations were chosen by one, two,
and one participant, respectively, while two participants excluded the end of the measurement resulting in strongly

Formatted: English (United States)

Formatted: English (United States)

Formatted: Heading 1, Indent: First line: 0 cm

negative, lower negative, stronger negative and intermediate CH_4 fluxes, respectively. Overall, the mean of the flux calculated by the 12 experts keeping this flux was $11.36 \text{ mgCH}_4 \text{ m}^{-2} \text{ d}^{-1}$ with a CV of 17%.

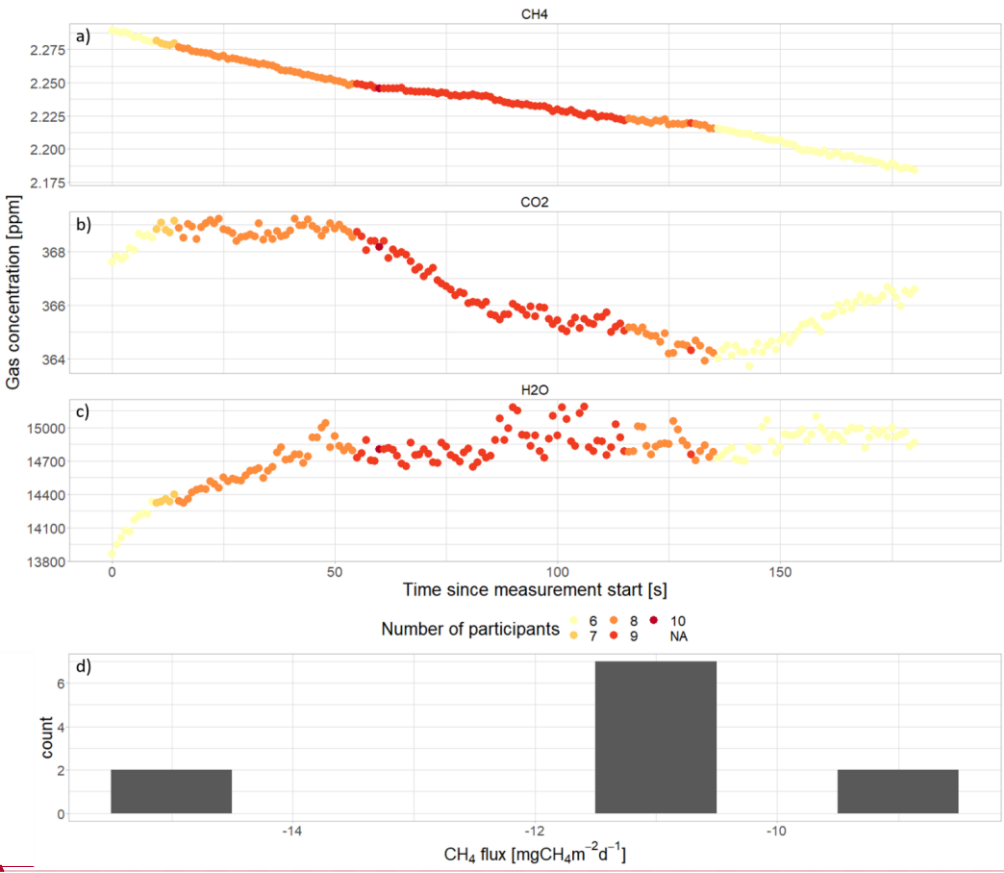


Figure 17: Measurement example VQC3 of a small linear decrease in CH_4 concentrations over the time of the chamber closure. Simultaneous measurements of CH_4 (a), CO_2 (b), and H_2O (c) concentrations over time during chamber closure with the colours of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses

Formatted: Font: Not Italic, English (United States)

Formatted: Heading 1

Formatted: Font: Not Bold, English (United States)

Formatted: Font: Not Bold, English (United States)

by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

3.2.8 No trend

In the visual QC exercise, we included one measurement example for which the CH₄ concentrations did not show a clear trend and varied only little over the time of the chamber closure (measurement ID VQC6, Figure 18). Most participants (69%) noticed the very small change in CH₄ concentrations over the whole measurement but they disagreed on whether the concentration measurements represented a real flux. Half of them suspected a real emission that remained very small because of CH₄ production and oxidation cancelling each other out at a low water table and two more participants called it a “zero flux” where the uncertainty would likely exceed the flux magnitude. Some (39%) of the participants, however, explained the low change in CH₄ concentrations by air leaking from the chamber, two of whom related the leak to vegetation obstructing the chamber seal and one to lateral diffusion into the chamber from the surrounding area.

Some (19%) of the participants furthermore pointed out an inconsistent trend in the CH₄ concentrations which they related to a changing balance between CH₄ production and oxidation over time, noisy measurements due to a low precision of the gas analyser, or a bad chamber seal combined with wind disturbance. According to one participant the latter was supported by the fluctuations appearing in the concentrations of all three gases, while two other participants mentioned that the CO₂ concentrations looked linear, at least after 30 to 40 s into the measurement, indicating an intact chamber seal.

The majority of the participants (62%) decided to discard the measurement due to leakage from the chamber (38%), a changing trend in the CH₄ concentrations (44%), a too short measurement time (13%), or too higher initial concentrations of CH₄ and CO₂ (13%). While two of these participants manually set the CH₄ flux to zero, one participant pointed out that the concentration changes were too large to be below the precision of the instrument so that the measurement should not be accepted as a zero flux. Some (31%) of the participants kept the measurement assuming a small but nonetheless real CH₄ flux and gave start and end times for flux calculation. Half of them discarded the beginning of the measurement as a period of initial equilibration, while the other half kept the entire measurement.

Formatted: English (United States)

Formatted: Heading 1, Indent: First line: 0 cm

Formatted: English (United States)

Formatted: English (United States)

The choice of different time periods for flux calculation by the participants resulted in a CV of 57% for this measurement example.

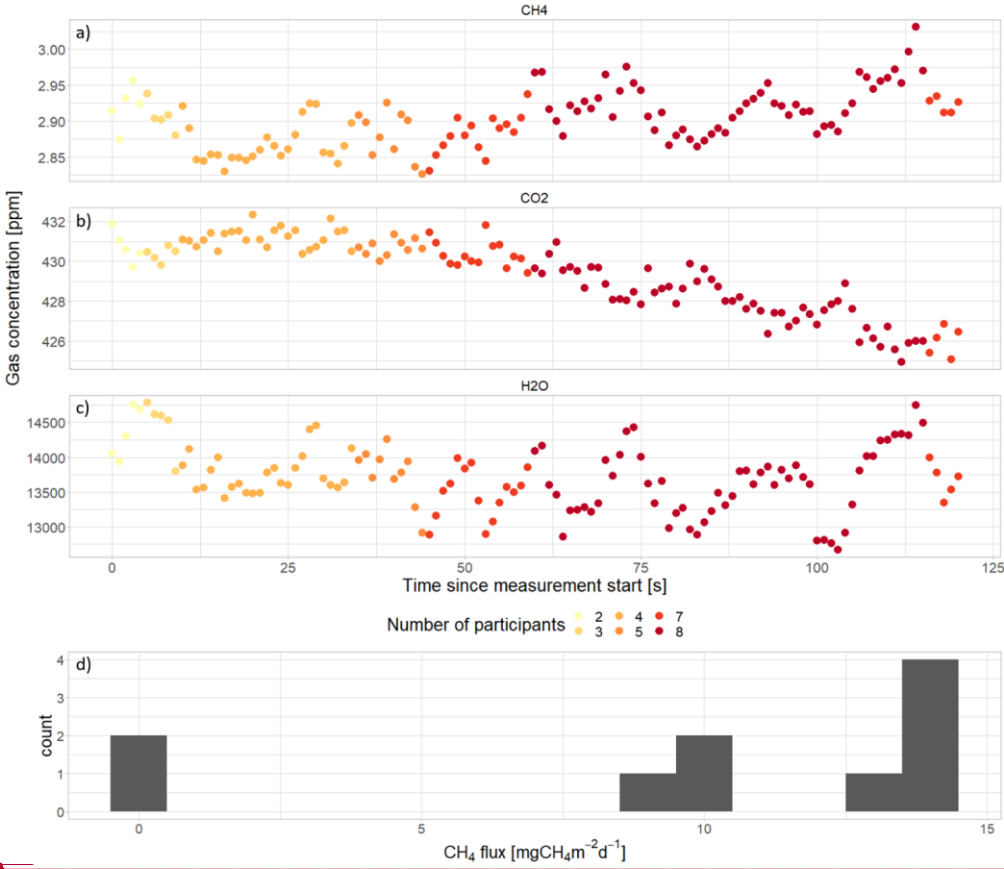


Figure 18: Measurement example VQC6 with the CH₄ concentrations showing little variation without a clear trend over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colours of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram

Formatted: Font: Not Italic, English (United States)

Formatted: Heading 1

Formatted: Font: Not Bold, English (United States)

Formatted: Font: Not Bold, English (United States)

of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants including the two participants who set the flux to zero (d).

Formatted: English (United States)

Formatted: Font: Not Bold, English (United States)

Formatted: English (United States)

3.2.29 Researcher variability and its implications for chamber flux synthesis studiesEffect of measurement period and QC on overall flux estimate using a representative dataset

Some of the measurement classes that we defined to represent different patterns in CH₄ concentrations over the time of the chamber closure clearly occurred more frequently than others within the total of 788 measurements that we took at Siikaneva bog (Table 1). The majority of the measurements (almost 60%) showed a linear increase in CH₄ concentrations in the chamber over the entire measurement. Nonlinear shapes with a decrease in slope over time were found in almost 20% of the measurements. Jumps in the CH₄ concentration either at the beginning or in the middle of the measurement were observed for 16% of the measurements while the remaining classes of an increasing slope over time, an inconsistent trend in CH₄ concentrations, linearly decreasing CH₄ concentrations over time, and no trend in CH₄ concentrations each made up less than 5% of the data set. Using the frequency of occurrence of each measurement class in the Siikaneva data set, we estimated an overall uncertainty of 28% introduced by different researchers discarding different numbers of measurements and an uncertainty of 17% due to different researchers choosing different parts of the same measurement for flux calculation.Overall, participants kept between 33-100% of the VOC flux examples for varying reasons although one participant discarded all fluxes.

Formatted: Heading 1, Indent: First line: 0 cm

Formatted: English (United States)

To estimate large-scale northern high-latitude CH₄ budgets, synthesis datasets of chamber CH₄ fluxes have been increasingly used (Kuhn et al., 2021; Treat et al., 2018). Generally, these synthesis datasets show consistent differences in CH₄ emissions among different wetland classes, but additionally, high variability within each wetland class that is attributed to high spatial and temporal variability and can be partly compensated for by using longer integration times (Treat et al., 2018) or by capturing spatial variability due to microtopography effects (Virkkala et al., 2024).

Formatted: Heading 1

Formatted: Subscript

Field Code Changed

Formatted: English (United States)

Formatted: Subscript

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

The results from this expert survey show that differences in methodology may be an additional factor contributing to high variability in CH₄ fluxes found in synthesis datasets. For example, the handling of low positive and negative fluxes can significantly affect estimates of CH₄ budgets, particularly in high-latitude and upland regions where low CH₄ emissions and/or uptake of CH₄ can be expected during large parts of the year. Discarding low fluxes or setting them to zero can therefore lead to a bias towards higher CH₄ emissions and potentially make the difference between a net annual uptake or a net emission of CH₄ in low-flux regions.

Formatted: Heading 1, Indent: First line: 0 cm

Formatted: Subscript

Formatted: Subscript

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Using the prevalence of different measurement scenarios in the Siikaneva data set (Table A1), we estimated an overall variability in flux estimates of 17% due to different researchers choosing different parts of the same measurement for flux calculation and a variation in the percentages of measurements passing the visual QC of 28% (Table A2). These estimates compare with the mean natural temporal variability of 19% but are lower than the mean natural spatial variability in CH₄ fluxes of 88% calculated from autochamber measurements in five temperate and Arctic peatlands by Pirk et al. (2016). Pirk et al. (2016) similarly found that both natural spatial and temporal variability in CH₄ fluxes exceed the difference between fluxes estimated using different fit functions. However, it has to be noted that the

185 uncertainty estimates derived in our study consider only the effect of differences in visual QC and do not account for different measurement setups and routines or different fit functions used for flux calculation. The entire uncertainty introduced into synthesis flux data sets by researcher variability might therefore exceed our estimates.

190 Our assessment of flux variability points towards the questions of where and when we introduce the largest error into our flux estimates — is it when we choose our measurement setup and processing approaches or do the location and the timing together with the spatial and temporal resolution of the measurements matter more? Answering this question will help identify the most relevant starting points for improving the accuracy of flux estimates and help lower uncertainties for flux syntheses. In any case, the survey shows that our human decision making introduces uncertainties that can obscure natural spatial and temporal variability in CH₄ fluxes. This might make it harder to identify relevant spatial and temporal environmental drivers of CH₄ fluxes, which is crucial for model development and CH₄ budget estimations.

195 4 Evaluation of the survey methodology

4.1 Expert survey approach and choice of question types insights

For our study, we decided to use the method of an expert survey which allowed us to combine the accuracy of a literature review with the directness of an expert assessment. In general, a literature review might provide a more complete overview of the methods used within a research community and thus allow for more reliable statistical interpretation of the results. However, we found that published data sets and research articles involving chamber fluxes are often lacking detailed information on measurement and data handling procedures – one of the current hurdles in interpreting, reusing, and combining existing chamber flux data sets. The expert survey, on the contrary, allowed us to obtain specific information directly from the source scientists and thereby collect data that might not be available in published literature but that might nonetheless significantly affect the CH₄ fluxes estimated from chamber measurements. In designing our survey, we had somewhat limited examples to follow as the approach of an expert survey rather than an expert assessment is not so commonly employed. While the exact implementation of the survey could therefore surely be refined in future studies, we showed here that surveying experts on their methods can be a useful approach that we can recommend as a beneficial line of research, and was strongly complementary to earlier reviews and recommendations of best measurement practices (e.g., Clough et al., 2020; de Klein and Harvey, 2012; Fiedler et al., 2022) **REFS**. The survey results clearly reveal that agreement on the measurement setup is high and generally in line with recommendations (Figure 4), but strong variability in the flux estimates (+18–28%) is introduced at the data processing and analysis stage by the different researchers (Figure 7, Table A2). This provides an opportunity to re-focus the discussion from measurement setups and linear vs. exponential fitting approaches to a wider discussion about data workflows and uncertainty sources in chamber flux measurements that have emerged with these new observational methods. In offering a diversity of question types, we ensured a balance between making the responses comparable and categorizable among the participants while still obtaining detailed information on their reasoning for the use of specific measurement and data handling techniques. The limited number of survey participants required a low number of possible responses in choice questions to allow for a meaningful statistical interpretation of the survey results. Yes/No questions further allowed us to draw

Formatted: Heading 1

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States), Subscript

Formatted: English (United States)

Formatted: Heading 2

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Comment Text, Indent: First line: 0,5 cm, Line spacing: 1,5 lines

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Subscript

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Field Code Changed

Formatted: English (United States)

Formatted: Highlight

Formatted: Font: 12 pt, English (United States)

Formatted: English (United States)

Formatted: Normal, Indent: First line: 0,5 cm

conclusions on the prevalence of the implementation of recommended best measurement practices among the survey participants.

4.2 Representativeness of the survey respondents and questions

From the variety of survey responses, it becomes clear that evaluating the representativeness of the respondents of the chamber flux community as a whole is challenging. One reason for this might be is that the chamber flux measurement community remains less organized than the eddy covariance flux measurement community and is more fluid, potentially because the barriers for entry are lower, i.e. the cost of analysis is lower. We recruited the survey participants from different places of employment assuming that this would make them rather independent in their choice of measurement and data handling approaches. Furthermore, we specifically reached out also to flux experts who are not represented in professional networks, expecting that a lower exchange with the rest of the chamber flux community might promote a stronger deviation from the commonly used methods. The main strength of the collected data set therefore lies in representing a large range of measurement and data handling practices; indeed, there were substantial and therefore demonstrating the maximum deviations in workflows applied within the part of the chamber flux community represented in this survey (Figures 4, 6). However, there might still be parts of the we did not reach all researchers using chamber fluxes -community that we did not reach with our survey; we likely or that are underrepresented in our study, for example those working in agricultural ecosystems, disturbed sites, and tropical ecosystems. The absence of the part of the chamber flux community that likely often observes low or close-to-zero CH₄ emissions or even CH₄ uptake. Overall, participants who had not encountered a certain shape in CH₄ concentrations in their own data sets before were more likely to discard the respective measurement example (Table 1, Table A2^{REF to Fig/table here}). For example, t_T might be reflected in the high discard rate in our visual QC exercise of measurementshe measurementexample, showing decreasing CH₄ concentrations over time was discarded by 50% of the current participants (Table 1), many of whom focused on wetland ecosystems (Table 1Figure 3), but is more likely to occur in well-drained agricultural soils (Mosier et al., 1997)(^{REF Mosier et al., 1997}). Overall, participants who had not encountered a certain shape in CH₄ concentrations in their own data sets before were more likely to discard the respective measurement example. This indicates that the handling of chamber measurements depends on the previous experiences of the researcher. As the prevalence of certain measurement scenarios depends on the ecosystem type researchedThus, the background of the survey participants might thus have affected the outcome of the visual QC exercise with a bias towards expected (higher) fluxes (Table 1). However, the handling of the example measurements can still be considered representative as the data set was collected in a boreal peatland—the ecosystem type that is most commonly researched by the survey participants (Figure 3). Additionally, the question of number of survey participants is always a concern. While the number of 46 researchers contacted (n=46) and the final maximum of 36 respondents might seem relatively low for a community survey, we estimate that this still represents a consideringconsiderable extent of the limitedan estimated total number of several hundred chamber flux experts world-wide. Time is always a factor in voluntary survey participation; therefore, it was important to streamline questions to incentivize survey completion. , the survey data set, besides demonstrating the range of methods used, also allows

Formatted: Font: 10 pt

Formatted: Heading 2

Formatted: Font: 10 pt

Formatted: Comment Text, Indent: First line: 0,5 cm, Line spacing: 1,5 lines

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt, Subscript

Formatted: Font: 10 pt

Formatted: Font: 10 pt, Subscript

Formatted: Font: 10 pt

Formatted: English (United States)

Formatted: Highlight

Formatted: Font: 10 pt

Formatted: Font: 10 pt

Formatted: Font: 10 pt, Subscript

Formatted: Font: 10 pt

Formatted: Highlight

Formatted: Font: 10 pt

Formatted: English (United States)

Formatted: Not Highlight

Formatted: English (United States)

for careful statistical interpretation, evaluating the prevalence of certain measurement, processing, and QC techniques. In offering a diversity of question types, we attempted a balance between making the responses comparable and categorizable among the participants while still obtaining detailed information on their reasoning for the use of specific measurement and data handling techniques. The limited number of survey participants required a low number of possible responses in choice questions to allow for a meaningful statistical interpretation of the survey results; therefore, we used yes/no answers rather than scales of agreement. Yes/No questions further allowed us to draw conclusions on the prevalence of the implementation of recommended best measurement practices among the survey participants.

4.3 Representativeness of the Siikaneva data set Assumptions in the flux calculations: site and researcher differences

Our estimates of researcher variability in flux data sets, derived from the visual QC exercise, strongly depended on the prevalence of different measurement scenarios in the underlying reference data set collected at Siikaneva Bog (Table A1). It is therefore crucial to evaluate the representativeness of this data set collected at Siikaneva bog. As the patterns in CH₄ concentrations represented in the different measurement scenarios are caused by both natural processes and chamber-induced artefacts, occur and their prevalence their occurrence depends on both the environmental conditions of the research site as well as on the chamber design and measurement setup.

Most measurements in the Siikaneva data set (~60%) showed the linear increase in CH₄ concentrations in the chamber headspace over time that is expected for an undisturbed measurement at a wetland site. However, a nonlinear, weakening increase in CH₄ concentrations was also represented by a rather high share of measurements (18%); and that Published studies confirm that this concentration pattern is also regularly observed in CH₄ concentrations from chamber measurements at other sites (e.g. Pirk et al., 2016). The survey responses confirm that it is often unclear whether this shape is caused by an initial disturbance of the measurement or by CH₄ saturation of the chamber headspace over time (Table 1). Furthermore, this lack of process-understanding shows through in the high variance associated with the non-linear fluxes (Table 1, Figure 7, Table A2). An initial disturbance, i.e. ebullition caused by the chamber placement, was a common explanation (Table 1) and might have occurred more frequently than average in the Siikaneva data set-, than other sites, as roughly 60% of the measurements were obtained from vegetation removal plots. The removal of vascular plants and of the *Sphagnum* moss layer might have reduced both plant-mediated CH₄ transport and CH₄ oxidation, resulting in higher CH₄ concentrations in the pore water and thus increasing the probability of ebullition events (Jentsch et al., 2024a). While CH₄ ebullition is a natural phenomenon often encountered in wetlands (Green and Baird, 2013), the increased probability of both natural and anthropogenically induced ebullition due to vegetation removal might have contributed to the high share of measurements (16xx%) showing abrupt jumps in CH₄ concentrations in the Siikaneva data set.

Although some measurement scenarios included in the visual QC exercise are relatively uncommon, it is still important to evaluate how these Evaluating how such some relatively uncommon a measurement scenarios would be handled by different researchers by including it in the visual QC exercise is still therefore still seems relevant important as theynd showed the largest

- Formatted: English (United States)
- Formatted: Indent: First line: 0,5 cm
- Formatted: English (United States), Not Highlight
- Formatted: English (United States)
- Formatted: Font: 10 pt, Not Bold, Not Italic, English (United States)
- Formatted: Heading 2, Space Before: 0 pt, After: 0 pt
- Formatted: Font: 10 pt
- Formatted: Font: 10 pt
- Formatted: Font: 10 pt
- Formatted: Font: 10 pt
- Formatted
- Formatted: Font: 10 pt
- Formatted: Font: 10 pt
- Formatted: Font: 10 pt
- Formatted: Font: 10 pt
- Formatted: Font: 10 pt
- Formatted: Subscript
- Formatted: Subscript
- Formatted: English (United States)
- Field Code Changed
- Formatted: English (United States)
- Formatted: Subscript
- Formatted: Highlight
- Formatted: English (United States)
- Formatted: Font: Italic
- Formatted: Font: Italic
- Formatted: Subscript
- Formatted: Subscript
- Formatted: Subscript
- Field Code Changed
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: Not Highlight
- Formatted: Highlight
- Formatted: Subscript
- Formatted: Not Highlight
- Formatted: Not Highlight
- Formatted: Not Highlight

researchers said they discarded from their own data sets was 5%, they discarded 19% the median share of measurements discarded in the visual QC exercise amounted to 19% when weighted by the considering the prevalence of the different measurement scenarios within the Siikanen data set (Tables A1, A2). Another reason for the high discard rate might be that related to the survey participants did not having done the example measurements themselves. While the median percentage of measurements that the researchers discarded from their own data sets was 5%, the median share of measurements discarded in the visual QC exercise amounted to 19% when considering the number of occurrences of the different measurement classes in the Siikanen data set. This implies that might be related to the They researchers did not have the option not being able to redo the measurements that they diagnosed as disturbed, in the visual QC data set as opposed to their own field measurements. Additionally, the Siikanen data set, from which the weighting factors for the different measurement classes included in the visual QC exercise were derived, might have contained more measurements that deviate from the expected continuous linear increase in CH_4 concentrations in the chamber than the data sets usually collected by the survey participants. As discussed earlier, this might be related both to the specific measurement setup used but also to the site-specific environmental conditions at Siikanen bog, and they lacked an overall view of the dataset. While the discard rate in the visual QC exercise might therefore not reflect how researchers handle their own field measurements, it might still occur that researchers need to process the chamber measurements collected by a colleague. Furthermore, for the purpose of assessing the uncertainty introduced into a flux data set by different data handling procedures a common data set had to be used. However, several participants mentioned that they would like to see the entire data set before deciding on keeping or discarding an individual measurement as they did not know the prevalence of the different measurement scenarios in the data set. Although required for reasons of practicability, the decisions for processing an entire data set might differ from the limited number of example measurements might render the results from the presented here visual QC exercise in our study not entirely representative of the decisions that researchers would make when processing an entire data set. Processing the full dataset as a common dataset rather than a small subset would also eliminate:

Presenting an entire data set for processing by the survey participants would furthermore remove the source of error introduced by our assumptions with the visual categorization classification of measurement scenarios (Table A1); however, this might also have decreased the number of respondents as this is a relatively intensive exercise. If respondents did their own flux calculations, this would allow for non-linear fitting methods, which we did not use in our exercise despite being occasionally suggested by a participant (7% of responses). While our this fitting and calculation approach may have been overly simplistic, post-hoc assumptions of how many participants would have used a non-linear fit and the different fitting options (such as exponential, quadratic, or logarithmic functions) would introduce substantial additional uncertainty into our the flux estimates of researcher variability. Reproducing the calculation approaches of every respondent would have required additional, very detailed information from the survey participants, likely reducing the number of completed surveys and making our uncertainty estimates less representative of the entire chamber flux community. However, this type of exercise might be worth undertaking in the future.

Formatted: English (United States)

As can be seen from the results of the visual QC part of the survey, the handling of different measurement examples assigned to the same class of measurement scenarios can differ strongly (Figure 7), even when processed by the same researcher. This indicates that our categorization might have been flawed, considering that similar patterns in CH₄ concentrations can arise from very different processes. However, the categorization of the data set into different measurement scenarios was highly beneficial in identifying major uncertainties in data handling and thus in deriving important starting points for future research.

4.4 Flux calculation in the visual QC exercise

For reasons of consistency, we decided to use a linear fit for flux calculation in the visual QC exercise even in the few cases where a participant suggested to use a nonlinear fit instead. This might have reduced the informative value of the variance in flux estimates for measurements showing a nonlinear change in CH₄ concentrations. However, we did not explicitly include the option of choosing between different fit functions in the visual QC exercise and can therefore not guarantee that all participants considered this possibility when selecting a time period for flux calculation and describing their handling of a measurement example. Furthermore, allowing for different fit functions would have added much complexity to our uncertainty assessment considering the variety in calculation methods mentioned by the survey participants as well as the different options for nonlinear fitting such as exponential, quadratic, or logarithmic functions. Exactly reproducing the calculation approaches of every respondent would have required additional and very detailed information from the survey participants. When setting up the survey, we therefore considered sharing a common data set of raw chamber measurements and asking the participants to process the data themselves, each using the flux processing and QC procedures that they typically use for their own measurements. While this might be a highly promising approach in the future, we decided against it for this study out of concern that the higher time effort required would further reduce the number of participants, making our uncertainty estimates less representative of the entire chamber flux community.

Estimating the effect of including the option of using a nonlinear fit on the researcher variability in flux estimates remains challenging: A linear fit leads to a lower flux estimate than the initial slope of a nonlinear fit in the case of a nonlinear increase in CH₄ concentrations with a decreasing slope but to a higher flux estimate in cases where the slope in CH₄ concentrations increases over time. Depending on the shape of the concentration change as well as on the time period chosen for flux calculation and the responses of the other participants using linear as opposed to nonlinear fits might have thus either increased or reduced the variance of the flux estimates.

5.4 Visions for improving the accuracy and compatibility of chamber CH₄ flux measurements and data sets

5.1 Recommendations for high-frequency measurements of CH₄ fluxes from chambers

Earlier studies have highlighted variability in CH₄ fluxes due to chamber design and fitting approaches (e.g., Fiedler et al., 2022; Maier et al., 2022; Pihlatie et al., 2013; Pirk et al., 2016; REF, Meier et al., 2022; Pihlatie et al., 2013). Here, we

Formatted: Strikethrough
Formatted: Indent: First line: 0,5 cm
Formatted: Strikethrough, Subscript
Formatted: Strikethrough

Formatted: Heading 2, Indent: First line: 0 cm
Formatted: Indent: First line: 0,5 cm

Formatted: English (United States), Strikethrough
Formatted: Strikethrough

Formatted: Indent: First line: 0,5 cm
Formatted: English (United States), Strikethrough
Formatted: Strikethrough
Formatted: English (United States), Strikethrough
Formatted: Strikethrough
Formatted: English (United States)
Formatted: English (United States), Subscript
Formatted: English (United States)
Formatted: Subscript
Formatted: Indent: First line: 0,5 cm
Formatted: Subscript
Field Code Changed
Formatted: German (Germany)
Formatted: German (Germany)

show that many researchers have adopted the recommended measurement techniques and setup (Figure 4). The relatively widespread adoption of high frequency CH₄ analyzers provides new challenges and illustrates a need to move focus from measurement setup and curve fitting ~~considerationaleculations~~ to data handling as the disagreement in QC approaches varies widely among the survey participants and explanations for some observed behaviors remain ~~inconsistent-muddled~~ (Tables 1, A2). While broader discussions about QC approaches are warranted, some simple steps may help to improve data quality:

- 1) **Calculation and implementation of a minimum detectable flux given the analyzer precision and chamber height.** Use this to determine the measurement length and to determine when fluxes are below detection limit. Use short measurement times to avoid chamber effects.
- 2) **Do not discard fluxes, including ebullition fluxes, low fluxes or zero fluxes.** Instead, we should move towards a ~~standardized QCquality control~~ flagging system. Ebullition fluxes, low and zero fluxes should be preserved and can be flagged in archived data. CO₂ concentrations can be used in addition to CH₄ concentrations to determine measurement quality (Pirk et al., 2016). This will work best in dark chambers as a net emission is expected. H₂O vapor is less reliable as an indicator of flux quality.
- 3) **Report all data forin archival purposes and implement data quality flagging.** A flagging system will indicate to others interested to re-use the data where uncertainties lie and has been implemented in eddy-covariance networks. Ideally, raw concentration data will be archived as well as processed data. This will allow reprocessing of data in the future as needed.

On the longer term, we need to develop new tools and networks to figure out how we can best ~~leverage the new possibilities~~ of high-frequency gas concentration measurementsuse all of this new data. Key steps are underway to allow easier operation, analysis, and standardization of flux calculation, for example the GoFlux Package for R (REF)(Rheault et al., 2024).- In earlier times, ebullition was difficult to identify using GC analysis but can be seen in the high frequency concentration time seriesfluxes (Fig. A1.6-A1.8), allowing the separation of ebullition from diffusive and plant-mediated fluxes (e.g., Hoffmann et al., 2017). The survey showed strong disagreement with how to handle these measurementsfluxes, sometimes resulting in quite large variations in flux magnitude (Figure 7), suggesting that this new insight into CH₄ transport pathways is not fully utilized. Overall, more discussion and exploration about this crucial measurement approach is needed to fully leverage the technological developments of the past decade.

5.2 Establish a formal trace gas chamber flux network

One reason for the large variability in chamber methods revealed in this survey could be a lack of exchange between the researchers working with chamber measurements of CH₄ fluxes. Only half of the survey participants are part of a flux monitoring network, such as FluxNet or AmeriFlux, both of which strongly focus on eddy covariance measurements. This indicates that the exchange within the chamber flux community might be impeded by a lack of suitable networking platforms. Chamber technique-focused conference sessions and workshops to further develop approaches and revise methodologies would be beneficial. Further discussion and recommendations toward a more rigorous standardization of flux calculations by

Formatted: Subscript

Formatted: List Paragraph, Numbered + Level: 1 +
Numbering Style: 1, 2, 3, ... + Start at: 1 + Alignment: Left
Aligned at: 0,63 cm + Indent at: 1,27 cm

Formatted: Font: Not Bold

Formatted: Font: Not Bold, Subscript

Formatted: Font: Not Bold

Formatted: Subscript

Formatted: English (United States)

Formatted: Subscript

Field Code Changed

Formatted: List Paragraph, Numbered + Level: 1 +
Numbering Style: 1, 2, 3, ... + Start at: 1 + Alignment: Left
Aligned at: 0,63 cm + Indent at: 1,27 cm

Formatted: Indent: First line: 0,5 cm

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: Subscript

Formatted: Not Highlight

1415 identifying the best fit based on objective criteria (e.g., Pedersen et al., 2010; Rheault et al., 2024) would be domain of such a
network. While much work has already gone into developing chamber-based approaches and recommendations for
measurements, the substantial (and potentially novel) uncertainty in fluxes calculated among researchers here indicates that
this matter is not yet settled (Figure 7). Furthermore, there was never complete agreement on whether to keep or exclude the
fluxes included in the survey (Table 1).

1420 Introducing a chamber flux network and data platform might speed adoption of a more standardized measurement protocol
(although many recommended chamber components are widely adopted, Figure 4), improve metadata and ancillary
measurements quality, spur development of a data quality flagging system that could foster a transparent exchange between
researchers on measurement and data handling procedures and ultimately enhance the compatibility of individual flux data
sets. Such a chamber flux network could build on existing research infrastructures such as the LTER sites, the ICOS sites in
1425 Europe and NEON sites in North America. Some examples for chamber databases have been developed like the Soil
Respiration Data Base (SRDB), which includes chamber measurements of ecosystem respiration and has been widely cited,
particularly for the response of ecosystem respiration to warming (B. Bond-Lamberty and Thomson, 2010; Ben Bond-
Lamberty and Thomson, 2010; Jian et al., 2020)(Bond-Lamberty & Thompson 2010a, 2010b, Jian et al., 2020). This requires
the open sharing of data, both raw chamber measurements and the calculated flux estimates using quality flagging rather than
1430 pre-filtered flux data sets. This way, reanalysis of existing data sets can be facilitated and all chamber measurements
contributing to larger scale synthesis studies can be reprocessed using a uniform calculation and QC approach to remove
differences among researchers, which we that we show have shown to makes a substantial difference in significantly affect
CH₄ flux estimates (Figure 7).

5.13 IntroducingDevelop and adopt a reference data settools for uncertainty estimation in data processing
1435 In our study, wWe demonstrated the potential of using a common data set to assess the varietyvariance in flux estimates
caused by different data processing and QC approaches (Figure 7). This approach could also be implemented more broadly to
build consensus on calculation methods, quality control, and data quality indices. We asked tThe survey participants to
performed a detailed, both quantitative and qualitative evaluation of few-example measurements, which allowed us to
comprehend and explained their decisions made in data handling (Figure 6). This helps to -and to thus identify the causes of
1440 deviations in differences and rationales for the processing and QC approaches they used; -Building on the knowledge gained
and having demonstrated the significant impact of data handling procedures on the resulting flux estimates in a rough
estimation, in a next step, this these differences affect fluxes even in the small number of example fluxes in this survey (Figure
7). The resultant variability in fluxesis could be -impact could be assesseded -in a more exact and purely quantitativetrigorous
way. For this, instead of just a few example measurements, by distributing an entire data set of raw chamber measurements
1445 could be shared with the flux experts, asking them and the community as a whole to process the data set using their own
calculation and QC approaches. Comparing the resulting flux valuesUsing a common dataset between the researchers could
provide us with a more accurate estimate of the potential uncertainty introduced into synthesis data sets due to data processing

Formatted: Not Highlight
Formatted: English (United States)
Formatted: English (United States)

Formatted: Normal, Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: Not Highlight

Formatted: Subscript

Formatted: Heading 2, Indent: First line: 0 cm

by different researchers and add insights into the relevance of this additional source of variability in CH₄ fluxes; through comparison with-in-addition-to-known the natural spatial and temporal variability in the data set. Sharing an entire data set; this would also eliminate some assumptions made in data processing in this survey (Section 4.3). The fluxes estimated from the raw data set could then be uploaded to a chamber flux network website (for an existing platform, such as GgitHub,) to add to a growing pool of flux data sets computed from the same chamber measurements by different researchers. A flagging metric could be added to indicate data quality, which would aid modelers and others in interpreting and understanding noisy observations, which are common in ecosystems with high spatial and temporal heterogeneity. This way, every interested researcher could assess how the flux estimates based on their processing techniques relate to those calculated by other flux experts and could help to build agreement on how to handle unclear/risky cases of non-linear concentration changes-fluxes.

Depending on the range of environmental and flux conditions represented in the reference data set, the variability in flux estimates arising from different processing approaches could be compared to the natural spatial and temporal variability in CH₄ fluxes as identified by the individual researchers. Thereby the meaningfulness of comparing and combining flux data sets computed by different researchers can be assessed. Such a comparison has previously been used (2016) to evaluate the effect of using linear vs. nonlinear regression for flux estimation by Pirk et al. (2016).

Besides the use in another expert survey, such a reference data set of raw chamber measurements could additionally be published online and made available for everyone to download and process. The fluxes estimated from the raw data set could then be uploaded to a website to add to a growing pool of flux data sets computed from the same chamber measurements by different researchers. This way, every interested researcher could assess how the flux estimates based on their processing techniques relate to those calculated by other flux experts.

Such a standardized test and training data set could be made even more valuable through including simulated inAdditional synthetic data could be an important addition to actual chamber measurements in the reference dataset and would clarify the processes resulting in strongly divergent flux estimates (Figure 7). A forward model could be developed to simulate the change in CH₄ concentrations in a chamber headspace as the real flux overlaid by a combination of chamber-induced artefacts in response to environmental conditions (Hutchinson and Livingston, 2001), Hutchinson and Livingston, 2001). This way, in addition to the fluxes computed by other researchers, researchers could compare their flux estimates to the known “real” flux underlying a simulated measurement to see if they succeed inis is possible to detecting it against background noise and artefacts.

Measurement simulations can –and– also help to understand how CH₄ (and covarying CO₂) concentrations change in response to commonly cited measurement issues that might result in non-linear fluxes (Table 1). The simulations can therefore answer the key questions that appeared in the visual QC exercise – Which part of a nonlinear measurement should be used for flux calculation? Can the linear part of a measurement following an ebullition event still be used for flux calculation? How should close-to-zero fluxes be identified and handled? Does a decrease in headspace CH₄ concentrations indicate actual CH₄ uptake? Pirk et al. (2016) demonstrated that Applying a nonlinear model for flux calculations can lead to an overestimation of CH₄ emissions if the nonlinear change in CH₄ concentrations was not, as assumed, caused by a change in the gas

Formatted: Subscript

Formatted: Comment Text, Indent: First line: 0,5 cm, Line spacing: 1,5 lines

Formatted: Subscript

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

Formatted: Subscript

Formatted: English (United States), Highlight

Formatted: Subscript

Formatted: Subscript

concentration gradient over time (Pirk et al., 2016). Improved process understanding will help to avoid introducing bias into flux data sets through unsubstantiated handling of non-linear measurements. The model simulations can answer the key questions that appeared in the visual QC exercise—Which part of a nonlinear measurement should be used for flux calculation? Can the linear part of a measurement following an ebullition event still be used for flux calculation? How should close-to-zero fluxes be identified and handled? Does a decrease in headspace CH₄ concentrations indicate actual CH₄ uptake?

Models and experimental studies to assess the effect of instrumentation on the flux estimates have previously been used to derive guidelines for certain aspects of measurement setups, such as chamber dimensions, pressure vents, and airtight seals (Christiansen et al., 2015, 2011; Hutchinson and Livingston, 2001; Pumpanen et al., 2004). Additionally, model-derived metrics can be used in post-hoc quality control, as was demonstrated for the minimum detectable flux (MDF) metric by Nickerson (2016). Such metrics will help identify a standardized set of required metadata on chamber setup and experimental design and ancillary measurements that should be taken alongside CH₄ fluxes in addition to the various variables currently recorded for the specific applications of the survey participants (Figure 5). For example, using a model with inputs of the air-soil CH₄ concentration gradient in the soil together with soil porosity and other soil properties can help to assess the potential effect of headspace saturation on the CH₄ flux (Pirk et al., 2016), which was commonly cited as problematic in this analysis (Table 1). Introducing model-derived such metrics will therefore help scientists to make more informed decisions in selecting a time period within a chamber measurement for flux calculation, choosing a fit function, and filtering the data set. From the metrics, and potentially derive diagnostics for that can be used for quality control flagging can be derived procedures that foster standardization of quality control procedures while factoring in site-specific conditions.

Applying a nonlinear model for flux calculations can lead to an overestimation of CH₄ emissions if the nonlinear change in CH₄ concentrations was not, as assumed, caused by a change in the gas concentration gradient over time (Pirk et al., 2016). Models and experimental studies to assess the effect of instrumentation on the flux estimates have previously been used to derive guidelines for certain aspects of measurement setups, such as chamber dimensions, pressure vents, and airtight seals (Christiansen et al., 2015, 2011; Hutchinson and Livingston, 2001; Pumpanen et al., 2004). Using a model with inputs of the air-soil CH₄ concentration gradient in the soil together with soil porosity and other soil properties can help to assess the potential effect of headspace saturation on the CH₄ flux (Pirk et al., 2016), which was commonly cited as problematic in this analysis (Table 1). Model-derived metrics can be used as diagnostics for post-hoc quality control, as was demonstrated for the Minimum Detectable Flux (MDF) metric by Nickerson (2016). The model simulations can answer the key questions that appeared in the visual QC exercise—Which part of a nonlinear measurement should be used for flux calculation? Can the linear part of a measurement following an ebullition event still be used for flux calculation? How should close-to-zero fluxes be identified and handled? Does a decrease in headspace CH₄ concentrations indicate actual CH₄ uptake? Introducing such metrics will therefore help scientists to make more informed decisions in selecting a time period within a chamber measurement for flux calculation, choosing

Formatted: Indent: First line: 0 cm

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

515 a fit function and filtering the data set and potentially derive diagnostics that can be used for quality control flagging procedures.

5.2 Developing advanced diagnostics for quality control

520 The survey results clearly reveal that variability in the flux estimates computed by different researcher is not so much introduced at the instrumentation stage, as agreement on the measurement setup is high (Figure 4), but rather at the data processing and analysis stage. Considerable uncertainty particularly showed regarding the identification of chamber-induced artefacts in the timeseries of CH₄ concentrations recorded in the chamber headspace and its distinction from natural processes involved in the CH₄ cycle. This disagreement resulted in a large range of flux estimates especially for measurements showing a nonlinear, weakening increase in CH₄ concentrations over time. 525 Considering the rather frequent occurrence of such concentration patterns, biases produced by different processing approaches for such measurement scenarios likely represent one of the largest remaining sources of divergence between flux data sets processed by different researchers. For example, Pirk et al. (2016) demonstrated the danger of unsubstantiated handling of certain measurement scenarios by showing that the use of a nonlinear model for flux calculation can lead to an overestimation of CH₄ emissions in cases where the nonlinear change in CH₄ concentrations was not, as assumed, caused by a change in the gas concentration gradient over time. Future research should therefore investigate under which conditions certain chamber-induced artefacts can be expected to significantly affect the CH₄ concentration timeseries recorded during a chamber measurement. This will allow for more informed decisions on the handling of chamber measurements. Besides its use to produce synthetic chamber measurements, a forward model that simulates the change in CH₄ concentrations observed during a chamber measurement could help to explore the potential effects of chamber artefacts on the flux estimates depending on measurement setup and environmental conditions. 535

540 Models and experimental studies to assess the effect of instrumentation on the flux estimates have previously been used to derive guidelines for certain aspects of measurement setups, such as chamber dimensions, pressure vents, and airtight seals (Christiansen et al., 2015, 2011; Hutchinson and Livingston, 2001; Pumpanen et al., 2004). Alternatively, metrics derived from such models can be used as diagnostics for post-hoc quality control, as was demonstrated for the Minimum Detectable Flux (MDF) metric by Nickerson (2016). Firstly, computing such metrics will allow a researcher to estimate the potential influence of a certain chamber artefact on their chamber measurements, given their specific measurement setup and site conditions. The model simulations will answer the key questions that appeared in the visual QC exercise – Which part of a nonlinear measurement should be used for flux calculation? Can the linear part of a measurement following an ebullition event still be used for flux calculation? How should close-to-zero fluxes be identified and handled? Does a decrease in headspace CH₄ concentrations indicate actual CH₄ uptake? Introducing such metrics will therefore help scientists to make more informed decisions in selecting a time period within a chamber measurement for flux calculation, choosing a fit function and filtering the data set. Secondly, from the metrics, diagnostics can be derived for the quality control of flux estimates. While most published scripts for flux processing that acknowledge nonlinear changes in gas concentrations aim for a rigorous standardization of flux calculations by identifying the best fit based on objective criteria (e.g., Pedersen et al., 2010; Rheault et al., 2024), implementing such 550

Formatted: Heading 1

Formatted: Font: Not Italic

Formatted: Highlight

Formatted: Subscript, Highlight

Formatted: Highlight

Formatted: Subscript, Highlight

Formatted: Highlight

Formatted: Subscript, Highlight

Formatted: Highlight

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: Subscript

Formatted: Subscript

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: Subscript

Formatted: Subscript

Formatted: Highlight

advanced process diagnostics could make flux processing more adaptable to individual field sites while maintaining a standardized calculation and quality flagging procedure.

Developing a forward model for a realistic simulation of chamber measurements will furthermore help to identify a standardized minimum set of variables that should be measured alongside the gas concentrations in the chamber headspace for a physically founded quality control of the flux estimates. For example, CH₄ concentration measurements in the peat pore air or water can reveal the concentration gradient in the soil which, together with soil porosity and other soil properties, can help to assess the potential effect of headspace saturation on the CH₄ flux (Pirk et al., 2016).

To test the effect of the measures the quantitative part of the survey could/should be repeated to see if we can agree better

5.3 Establishing a chamber flux network

One reason for the large variability in chamber methods revealed in this survey could be a lack of exchange between the researchers working with chamber measurements of CH₄ fluxes. Only half of the survey participants are part of a flux monitoring network, such as FluxNet or AmeriFlux, most of which strongly focus on eddy covariance measurements. This indicates that the exchange within the chamber flux community might be impeded by a lack of suitable networking platforms. Introducing a chamber flux network and data platform involving a standardized measurement protocol, requirements for metadata and ancillary measurements, and quality flagging system could foster a transparent exchange between researchers on measurement and data handling procedures and enhance the compatibility of individual flux data sets. Such a chamber flux networks could build on existing research infrastructures such as the LTER sites, the ICOS sites in Europe and NEON sites in North America.

Furthermore, such a data base should promote the sharing of raw chamber measurements in addition to the flux estimates as well as quality flagged rather than pre-filtered flux data sets. This way, reanalysis of existing data sets can be facilitated and all chamber measurements contributing to larger scale synthesis studies can be reprocessed using a uniform calculation and QC approach.

6. Conclusions

Chamber flux measurements are crucial for capturing spatial variability in ecosystems and quantifying treatment effects on greenhouse gas emissions. At broader spatial scales, synthesis datasets of chamber CH₄ fluxes show differences among but also high variability within wetland classes (Kuhn et al., 2021; Treat et al., 2018). The high variability within classes has been attributed to high spatial and temporal variability and can be partly compensated for by using longer integration times (Treat et al., 2018) or by capturing spatial variability due to microtopography effects (Virkkala et al., 2024). The results from this expert survey show that differences in methodology may be an additional factor contributing to high variability in CH₄ fluxes across sites and datasets. For example, the handling of low positive and negative fluxes can significantly affect estimates of CH₄ budgets, particularly in high latitude and upland regions where low CH₄ emissions and/or uptake of CH₄ can be expected during large parts of the year. Discarding low fluxes or zero fluxes can lead to a bias towards higher CH₄ emissions (e.g. Table X) and potentially make the difference between a net annual uptake or a net emission of CH₄ in low-flux regions. Ebullition

Formatted: Subscript

Formatted: Subscript

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: Heading 1, Indent: First line: 0 cm

Formatted: Subscript

Formatted: English (United States)

Formatted: Heading 1

Formatted: Highlight

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: Not Highlight

Formatted: Not Highlight

Formatted: English (United States)

Formatted: Not Highlight

Formatted: English (United States)

585 events may also comprise a substantial fraction of emissions; discarding these may lead to an underestimation of ecosystem
CH₄ fluxes.

Our assessment of flux variability points towards the questions of where and when we introduce the largest error into our
flux estimates – is it when we choose our measurement setup and processing approaches or do the location and the timing
together with the spatial and temporal resolution of the measurements matter more? Answering this question will help identify
590 the most relevant starting points for improving the accuracy of flux estimates and help lower uncertainties for flux syntheses.
In any case, the survey shows that our human decision making introduces uncertainties that can obscure natural spatial and
temporal variability in CH₄ fluxes. This might make it harder to identify relevant spatial and temporal environmental drivers
of CH₄ fluxes, which is crucial for model development and CH₄ budget estimations.

595 The assumptions and the uncertainty among the survey participants surrounding their selection of approaches for flux
calculation and QC call for better diagnostics to unambiguously identify the processes that cause the observed patterns in the
CH₄ concentrations. The survey participants suggested various reasons for a nonlinear behavior in CH₄ concentrations (Table
2) which strongly affected their decisions during flux calculation and QC and led to a large variation in the resulting flux
estimates (Table 1, Figure 6). This poses the question of how to identify the actual reason for the observed nonlinearity which
needs to be known in order to make an informed decision on when to redo or to discard a measurement and how to handle a
measurement in flux calculation and QC in order to get an accurate estimate of the real CH₄ flux.

The occurrence for example of nonlinear changes in CH₄ concentrations or ebullition events might be influenced by the
measurement setup but likely also depends on the environmental conditions. This indicates that in order to derive
605 recommendations on how to best estimate the CH₄ flux from a given chamber measurement the environmental conditions
under which certain phenomena such as different types of chamber artefacts like leakage and saturation or ebullition events
can significantly affect the CH₄ concentrations in the chamber need to be identified. For example, as Pirk et al. (2016) showed,
the unsubstantiated use of a nonlinear model for flux calculation can lead to an overestimation of CH₄ emissions and should
therefore only be applied in cases where we can be sure that the nonlinear change in CH₄ concentrations was caused by a
610 change in the gas concentration gradient. Improved diagnostics to identify the processes involved will therefore allow for more
informed decisions on how to handle chamber measurements that show a nonlinear change in CH₄ concentrations and will
ultimately result in more accurate flux estimates. While most published scripts that acknowledge nonlinear changes in gas
concentrations aim for a standardization of flux calculations by identifying the best fit based on objective criteria (e.g., Pedersen
et al., 2010; Rheault et al., 2024), implementing such improved process diagnostics could make flux processing more adaptable
to individual field sites while maintaining a standardized calculation procedure.

615 Developing improved process diagnostics will also help to identify a standardized minimum set of variables that
should be measured alongside the gas concentrations for additional quality control of the flux estimates. The visual QC exercise
in this survey should be considered as an example of how researchers would handle a given data set rather than their own

Formatted: Subscript

Formatted: English (United States)

Formatted: English (United States)

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

1620 measurements as the additional information given in the visual QC exercise might not be entirely representative of the
additional variables that researchers would select to assess the quality of their flux estimates. For example, not all researchers
might usually consider the behavior of the CO₂ and H₂O concentrations when evaluating a CH₄ flux. In the visual QC exercise,
however, most participants took the other gas concentrations into account as they were available while one participant
mentioned that an unexpected pattern in the H₂O concentrations alone would not make them discard a CH₄ flux measurement.
While recommendations on the chamber setup and measurement routines are outlined in many places and well founded by
1625 modelling and experimental studies, the steps for data processing, including data fitting methodologies and QC approaches
might also have a strong but less well understood effect on the calculated CH₄ fluxes.
Only one third of the survey participants assessed the uncertainty of their individual flux estimates which similar to the MDF
allows for more profound quality flagging of the data. One participant reports the differences between flux values resulting
from different fit functions as uncertainty estimates. This allows to circumvent the decision between linear or nonlinear fit for
1630 flux calculation in cases where the reason for the nonlinearity in the CH₄ concentrations are unknown and it is thus not clear
which function better represents the real flux thereby preventing strongly biased data sets. Assessing the uncertainty of flux
estimates supports the request by Maier et al. (2022) that instead of being discarded all fluxes should be published in data bases
together with quality flags or uncertainty estimates.

Formatted: English (United States)
Formatted: English (United States)
Formatted: English (United States)

Formatted: English (United States)
Formatted: Indent: First line: 0 cm

Formatted: English (United States)

Formatted: English (United States)
Formatted: Normal

1635 **4.1 Implementation and development of Best Measurement Practices for chambers**

Numerous attempts have been made towards standardizing flux measurements using static chambers by compiling and publishing guidelines on best measurement practices on chamber measurements (e.g. de Klein and Harvey, 2012; Fiedler et al., 2022). Our expert survey revealed that despite the existing guidelines, not all researchers are implementing the recommended measures for various reasons (Figure 4, Table 3). Reasons for not implementing certain chamber equipment or processing advice could be that some researchers might simply be unaware of the recent guidelines on chamber measurements due to a lack of collaboration and a lack of suitable networking platforms. Only half of the survey participants are part of a flux monitoring network, such as FluxNet or AmeriFlux, most of which strongly focus on eddy covariance measurements. Other potential reasons for not implementing recommended measures include that existing equipment might not be suitable for adjustments, e.g. specific chamber sizes or shapes can make some measures inappropriate or impractical, financial constraints might not allow for improvements or new equipment, or there might be site specific requirements on the chamber design. The various scientific backgrounds of the researchers doing chamber measurements (Figure 2b) might further be contributing to the variety of measurement setups, calculation and QC procedures found in this survey as the educational training likely influences which aspects of the flux measurements, i.e. which chamber artefacts and environmental controls on CH₄ fluxes, are considered most important.

Field Code Changed
Formatted: English (United States)

1650 Some shortcomings in measurement setups can likely be related to an uncertainty around the effect of certain chamber equipment on the measured CH₄ flux, leading to concerns that overcompensating for certain chamber artefacts can introduce

Formatted: English (United States)

new sources of error. While most researchers use fans to mix the air inside their chamber, some researchers mentioned that their chambers are small enough to not need a fan. This statement highlights that further research is needed to investigate the strength of turbulence that is adequate for particular chamber dimensions to ensure proper mixing of the chamber air while preventing that additional gas is artificially released from the soil (Christiansen et al., 2011; Maier et al., 2022). Vents for pressure equilibration are likely only used by half of the participants for the similar fear of causing a Venturi effect when wind passes over the vent outlet (Bain et al., 2005; Conen and Smith, 1998). Clear guidelines exist however on how to avoid the Venturi effect by adjusting the vent design (Xu et al., 2006). Furthermore, the two types of vents—the one that is open only during chamber placement and the one that remains open during the measurement—seem to be considered rather as alternatives for vent designs than as two measures that tackle different pressure-related chamber artefacts and that should therefore both be applied simultaneously. The survey participants generally avoid the danger of overcompensating for a temperature increase inside the chamber and causing condensation (Fiedler et al., 2022) by using active cooling only in cases where it is required, e.g. because transparent chambers are used. Depending on the environmental conditions, opaque chambers could be used more often for insulation but blocking out the incoming radiation could reduce active CH₄ transport through plant aerenchyma thereby reducing the measured CH₄ emissions (Clough et al., 2020).

The ability to detect or correct for any remaining temperature and/or pressure differences between chamber and ambient air varies among the participants as not all of them record the temperature and/or pressure inside their chamber. Only two participants can account for temperature and/or pressure changes over the time of the chamber closure by individually correcting each concentration measurement as they document the chamber temperature and/or pressure at the same frequency as the gas concentrations. Changing temperature and pressure conditions inside the chamber might go unnoticed when using only one temperature and pressure reading to correct the final flux estimate. Most notably, 17% of the survey participants do not measure the chamber temperature at all (Figure 4), which can lead to large uncertainties considering the large linear effect of temperature on the flux magnitude through the ideal gas law.

Similar to controlling the temperature inside the chamber the requirements for chamber dimensions, chamber insertion depths into the soil, and chamber seals strongly depend on the environmental conditions of the research site. Chamber dimensions need to be adapted to the surface structure as well as to the vegetation height while allowing for flux detection within reasonable deployment times. As the required insertion depth of the chamber into the soil as well as the necessity of a gastight seal are low under water saturated conditions and at low soil porosities, the fact that one third of the survey participants did not use a collar or a seal on their chamber might be less problematic than it appears since many participants measure in wetlands or on open water. The chamber setup should nevertheless be tested for gas tightness before it is deployed in the field.

An increasing use of inline gas analyzers can loosen the requirements on chamber dimensions for CH₄ flux measurements due to shorter deployment times. In most researched environments CH₄ emissions are high enough so that the minimum detectable flux is reached quickly after the chamber closure. The high measurement frequency of inline gas analyzers therefore allows to significantly reduce the chamber deployment time, thereby decreasing the potential effect of chamber artefacts on the CH₄ flux estimates. Chambers can therefore be smaller as shorter measurement times reduce the potential

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

690

695

1700

effects of leakage, lateral diffusion, temperature and pressure changes on the flux estimates. Currently, the majority of the survey participants (80%) use inline gas analyzers. Of the 26% of participants who manually sample the chamber air, two participants (22%) keep their chamber closed for more than 40 min which is considered as too long by Rochette and Eriksen-Hamel (2008) but earlier guidelines allowed for up to 1 hour closure time (Holland et al., 1999). In addition to increasing the relevance of chamber artefacts due to longer closure times, manual sampling can obscure the influence of chamber artefacts through the lower temporal resolution at which the gas concentrations inside the chamber are monitored. This limits the possibilities to still exclude measurement periods affected by chamber artefacts at the stage of flux processing and quality control.

The different variables measured alongside the fluxes might indicate that depending on their background and research questions the survey participants consider different variables as important in controlling CH₄ fluxes. The ancillary variables also determine which additional information is available to the researchers to evaluate the quality of the CH₄ flux measurements. Almost all survey participants measured variables to characterize the soil, hydrological and meteorological conditions, covering most of the ancillary data suggested by Maier et al. (2022). The potential effects of the vegetation cover were however considered by less than one sixth of the participants only.

Table 3: Recommendations for chamber design (adapted from Clough et al., 2020) (Roman), their implementations by researchers as derived from the expert survey and resulting issues (bold).

Design feature	Design objective	Minimum requirements	Site-specific issues	Implementation by researchers	Evolving issues	Improvements
AREA	Minimize error due to poor sealing and maximize area sampled.	A chamber area/perimeter ratio of ≥ 10 cm is recommended. (equates to a cylindrical chamber of ≥ 40 cm diameter).	Adaption needed if rocks or roots are present, or if required by research objectives.	66% of chambers have area/perimeter ratio of ≥ 10 cm. 75% of chambers with area/perimeter ratio < 10 cm are cylindrical.		When using cylindrical chambers, make sure that they have a diameter of ≥ 40 cm.
HEIGHT	Maximize flux detection and minimize perturbation of	Chamber height (cm) to deployment time (h) ratio should be ≥ 40 cm h ⁻¹ .	Chamber height should accommodate crop height.	93% of measurement setups had chamber height to deployment time ratio of ≥ 40		Inline gas analyzers allow for shorter deployment times, so lower chamber heights

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: Heading 2

Field Code Changed

Formatted: English (United States)

	environmental variables.			cm h ⁻¹ . The two remaining setups had too long closure times considering the relatively low chambers.		are acceptable as minimum detectable CH ₄ flux is reached after short deployment in most researched environments.
BASE DEPTH	Prevent below ground lateral gas transport, shading and ponding of water.	Ratio of insertion depth: to deployment time of ≥12 cm h ⁻¹ . Height above soil surface should be as close to the soil surface as practical (<5 cm).	Required insertion depth is higher at lower soil porosity.	66% of the chamber setups involved a base (collar) inserted into the soil; 25% of the participants do aquatic measurements where no collar is needed, and 83% of the survey participants measure CH ₄ fluxes from wetlands, where saturated soil conditions allow for low insertion depths.	Lateral diffusion effects on the concentration increase?	Inline gas analyzers allow for shorter deployment times as minimum detectable CH ₄ flux is reached after short deployment in most researched environments, making shallower insertion depths are acceptable
GASTIGHT SEAL	Prevent gas leaking between chamber and base.	A water trough or rubber/closed cell foam gasket. Gaskets should have low internal cross-sectional	Appropriate sealing method depends on the environment. No sealing is required for aquatic measurements.	63% of the participants used a water trough or gasket to seal between chamber and base.	Many chamber setups do not use a seal despite clear recommendation. Leaky seal effects on concentration increases?	Chamber setup should be tested for gas tightness.

Formatted: English (United States)

Formatted: English (United States)

		area—and—be compressible; appropriate fasteners—are required—with rubber gaskets.		25%—of—the participants—do aquatic measurement where no seal is required.		
VENT			Tube-type	51% of chamber	Not used enough	Both vent type i)
i)——while placing chamber——on base	i) To prevent pressure disturbance while placing the——chamber on the base.	i) Opening a vent——or sampling——port while placing the chamber is essential.	vents need to be adapted to expected wind speeds to avoid Venturi effect.	setups involve some kind of vent. Responses indicate that both types of vents are rarely used simultaneously.	despite——clear recommendation.	and ii) should be applied. Danger of Venturi effect can be avoided following well-founded recommendation on the vent design.
ii)——during deployment	ii) To prevent pressure gradients between the interior——and exterior of the chambers during flux measurement and——gas sampling.	ii) Tube-type vents should be located close to the soil surface; or be designed to——minimize wind——effects. Appropriate vent dimensions (diameter——and length)——are dependent——on expected wind speeds during deployment and should be adjusted accordingly. Chambers and their——vents should——be bench tested to				

Formatted: English (United States)

		ensure no Venturi effect occurs. Designs exist to overcome Venturi effects.				
INSULATION	Prevent temperature gradients between the interior and exterior of the chambers.	Use reflective foil, foam, or polystyrene. Test effectiveness by comparing surface soil temperatures inside and outside the chambers	Chamber needs to be transparent if CO ₂ uptake through photosynthesis measured alongside the CH ₄ flux.	3% of the participants insulate their chamber and 17% actively cool the chamber air.	Cooling of the headspace air relative to the ambient air which might lead to condensation inside the chamber or sampling tubes (Fiedler et al., 2022).	Use cooling system only if chamber cannot be insulated and/or if long chamber deployment periods are needed (Maier et al., 2022).
HEADSPACE MIXING	Well-mixed headspace to ensure that representative sample is taken.	Active headspace mixing (e.g., fans) should not affect the diffusive flux.	Crop type and chamber height.	80% of the chamber setups involve fans. 43% of the participants who do not use a fan stated that the air flow from circulation through a closed loop with the gas analyser sufficiently mixes the chamber air, particularly in small chambers.	Missing concrete knowledge on how much mixing is adequate for which chamber dimensions.	Effects of mixing should be tested and reported on. There has been relatively little work performed on evaluating specific requirements for given chamber geometries and fan size wind speed combinations.

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

4.2 Considerations for data processing

While recommendations on the chamber setup and measurement routines are outlined in many places and well founded by modelling and experimental studies, the steps for data processing, including data fitting methodologies and QC approaches might also have a strong but less well understood effect on the calculated CH₄ fluxes.

A subjective assessment of different chamber effects led to the choice of different time periods for flux calculation and had the largest effect for measurements that showed a nonlinear increase in CH₄ concentrations (Table 1, Figure 6). Nonlinear measurements with a decreasing slope over time accounted for over 18% of our example data set, making it the second largest measurement class after linearly increasing CH₄ concentrations. Published studies confirm that this shape is regularly observed in CH₄ concentrations from chamber measurements (e.g. Pirk et al., 2016). Biases produced by different processing approaches for these measurements therefore likely represent one of the largest sources of divergence between flux data sets processed by different researchers. To avoid any initial disturbance caused by the chamber placement from influencing the flux estimate, almost half of the survey participants generally exclude the beginning of each measurement from their flux calculation. For measurements with a nonlinear increase in CH₄ concentrations levelling off over time, this removes the part of the measurement with the steepest increase in CH₄ concentrations resulting in lower flux estimates. In contrast, believing that the beginning of the measurement is least affected by chamber artifacts such as saturation, condensation, leakage, and temperature and pressure deviations from ambient conditions, about one third of the participants consider an exponential fit for their flux calculations, using the steepest, initial slope of the CH₄ concentrations as their flux estimate. This overall uncertainty on which part of a nonlinear measurement best represents the real flux also shows in the large range of flux estimates derived for the nonlinear measurements with decreasing slope in the visual QC exercise (Figures 6, 11). The visual QC exercise furthermore revealed that in extreme cases where the trend in CH₄ emissions reverses from an increase to a decrease over the time of the measurement, interpretations of the concentration time series make the difference between quantifying net CH₄ emission or uptake from the same measurement (Figures 6, 16).

The survey participants were also divided on the question of keeping chamber measurements that similarly show a nonlinear increase in CH₄ concentrations but with an increasing slope over time. This shape in CH₄ concentrations occurred less often in only 3% of the measurements in our example data set but was still reported surprisingly often also in other studies considering this behavior is not consistent with diffusion theory (Kutzbach et al., 2007). Accordingly, half of the participants who discarded the measurement supported their decision by stating that they cannot explain the observed shape in CH₄ concentrations and therefore cannot justify choosing a time period for flux calculation as this would largely affect the flux estimate considering the nonlinearity. The flux estimates computed for the 65% of participants who kept the measurement and gave start and end times for flux calculation confirm that this concern was justified as the flux estimates differ by up to 78% depending on the time period chosen.

The informative value of the variance in flux estimates for measurements showing a nonlinear change in CH₄ concentrations might however be reduced as, for reasons of consistency, we used a linear fit for flux calculation even in cases

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

where participants suggested to use a nonlinear fit. A linear fit leads to a lower flux estimate than the initial slope of a nonlinear fit in the case of a nonlinear increase in CH_4 concentrations with a decreasing slope but to a higher flux estimate in cases where the slope in CH_4 concentrations increases over time. Depending on the shape of the concentration change as well as on the time period chosen for flux calculation and the responses of the other participants using linear as opposed to nonlinear fits might have thus either increased or reduced the variance of the flux estimates.

The examples showing a nonlinear increase in CH_4 concentrations furthermore revealed that the QC procedure is subjective in the way that it depends on previous experiences. For example, participants who had not encountered a certain shape in CH_4 concentrations in their own data sets before were more likely to discard the respective measurement example in the visual QC exercise. How frequently certain types of shapes in CH_4 concentrations appear in a data set in turn likely depends on the ecosystem type that is researched. Several participants therefore mentioned that they would like to see the entire data set before deciding on keeping or discarding an individual measurement. Due to the limited number of example measurements, the visual QC exercise in our study might therefore not entirely represent the decisions that researchers would actually make when processing an entire data set.

CH_4 ebullition is another phenomenon often encountered in wetlands (Green and Baird, 2013), affecting more than 16% of the measurements in our example data set, and that is handled differently by different researchers thereby introducing additional uncertainty into data sets combining flux estimates from different researchers. Most importantly, the survey participants disagreed on whether ebullition events should be included in flux estimates from chamber measurements or if diffusive and ebullitive flux should be quantified separately, either by isolating periods of ebullitive and diffusive flux from one concentration time series or by separately measuring ebullition (e.g. Hoffmann et al., 2017), for example using bubble traps (e.g. Männistö et al., 2019). The visual QC of measurement examples showing jumps in the CH_4 concentrations furthermore revealed that there was some uncertainty surrounding the distinction between ebullition events and artefacts of the gas analyzer among the survey participants. The flux estimates of the 4% to 8% of the participants who would use a linear fit over an entire measurement containing episodic ebullition events to account for both diffusive and ebullitive CH_4 emissions were up to five times as high as the flux estimates from the participants considering the diffusive flux only (Figures 12, 13, 14). This highlights the importance of reporting whether ebullitive emission was included in a given flux data set.

The survey participants furthermore disagreed on whether the remaining part of a measurement after an ebullition event could still be used to quantify the diffusive flux. More than half of the survey participants (54%) kept the linear part of a measurement after an initial ebullition event for flux calculation while 38% of the participants discarded the entire measurement assuming that the high CH_4 concentrations in the chamber following the ebullition event would decrease the concentration gradient and thus decrease the CH_4 flux between soil and chamber headspace for the rest of the measurement (Figure 13). This effect also influenced the range of flux estimates from a measurement with repeated ebullition events (Figure 14). Flux estimates from the 15% of participants using the shorter linear increase in CH_4 concentrations before the first ebullition event were three times as high as the flux estimates from the 19% of participants fitting the longer linear increase after the first ebullition event for their flux calculations.

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Field Code Changed

Formatted: Subscript

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Similar to ebullition events, the conditions, and in particular the water table depths, under which net uptake of CH_4 can occur are debated within the flux community. Due to the linear development, flux estimates for a measurement showing a constant decrease in CH_4 concentrations over time differed less between the participants compared to nonlinear examples. However, more participants (54% compared to 12% to 31% for nonlinear measurement with constant trend) decided to discard the entire measurement because they do not expect a net uptake of CH_4 in a wetland despite the relatively low water table.

Another source of uncertainty lies in the identification and the handling of “zero fluxes”. Two thirds of the survey participants discarded a measurement example in the visual QC exercise showing only very low variations in CH_4 concentrations without a clear trend over the time of the chamber closure. The other third of the participants made a flux estimate, 20% of whom set the flux to zero and 80% calculated a small positive flux. Only one participant remarked that the magnitude in CH_4 variations needs to be compared to the instrument precision to decide whether a measurement can be classified as a “zero flux”. This approach gets closest to computing the minimum detectable flux (MDF) introduced by Christiansen et al. (2015) and refined by Nickerson (2016) to assess the quality of low flux estimates. Contrary to the approaches of most survey participants, Maier et al. (2022) recommend to not discard or set to zero fluxes estimates below the MDF.

Only one third of the survey participants assessed the uncertainty of their individual flux estimates which similar to the MDF allows for more profound quality flagging of the data. One participant reports the differences between flux values resulting from different fit functions as uncertainty estimates. This allows to circumvent the decision between linear or nonlinear fit for flux calculation in cases where the reason for the nonlinearity in the CH_4 concentrations are unknown and it is thus not clear which function better represents the real flux thereby preventing strongly biased data sets. Assessing the uncertainty of flux estimates supports the request by Maier et al. (2022) that instead of being discarded all fluxes should be published in data bases together with quality flags or uncertainty estimates.

4.3 Remaining knowledge gaps and next steps

The assumptions and the uncertainty among the survey participants surrounding their selection of approaches for flux calculation and QC call for better diagnostics to unambiguously identify the processes that cause the observed patterns in the CH_4 concentrations. The survey participants suggested various reasons for a nonlinear behavior in CH_4 concentrations (Table 2) which strongly affected their decisions during flux calculation and QC and led to a large variation in the resulting flux estimates (Table 1, Figure 6). This poses the question of how to identify the actual reason for the observed nonlinearity which needs to be known in order to make an informed decision on when to redo or to discard a measurement and how to handle a measurement in flux calculation and QC in order to get an accurate estimate of the real CH_4 flux.

The occurrence for example of nonlinear changes in CH_4 concentrations or ebullition events might be influenced by the measurement setup but likely also depends on the environmental conditions. This indicates that in order to derive recommendations on how to best estimate the CH_4 flux from a given chamber measurement the environmental conditions under which certain phenomena such as different types of chamber artefacts like leakage and saturation or ebullition events

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: Indent: First line: 0 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

can significantly affect the CH_4 concentrations in the chamber need to be identified. For example, as Pirk et al. (2016) showed, the unsubstantiated use of a nonlinear model for flux calculation can lead to an overestimation of CH_4 emissions and should therefore only be applied in cases where we can be sure that the nonlinear change in CH_4 concentrations was caused by a change in the gas concentration gradient. Improved diagnostics to identify the processes involved will therefore allow for more informed decisions on how to handle chamber measurements that show a nonlinear change in CH_4 concentrations and will ultimately result in more accurate flux estimates. While most published scripts that acknowledge nonlinear changes in gas concentrations aim for a standardization of flux calculations by identifying the best fit based on objective criteria (e.g., Pedersen et al., 2010; Rheault et al., 2024), implementing such improved process diagnostics could make flux processing more adaptable to individual field sites while maintaining a standardized calculation procedure. Developing improved process diagnostics will also help to identify a standardized minimum set of variables that should be measured alongside the gas concentrations for additional quality control of the flux estimates. The visual QC exercise in this survey should be considered as an example of how researchers would handle a given data set rather than their own measurements as the additional information given in the visual QC exercise might not be entirely representative of the additional variables that researchers would select to assess the quality of their flux estimates. For example, not all researchers might usually consider the behavior of the CO_2 and H_2O concentrations when evaluating a CH_4 flux. In the visual QC exercise, however, most participants took the other gas concentrations into account as they were available while one participant mentioned that an unexpected pattern in the H_2O concentrations alone would not make them discard a CH_4 flux measurement.

Similarly, the high number of measurements discarded in the visual QC exercise might be related to the survey participants not having done the example measurements themselves. While the median percentage of measurements that the researchers discarded from their own data sets was 5%, the median share of measurements discarded in the visual QC exercise amounted to 19% when considering the number of occurrences of the different measurement classes in the Siikaneva data set. This might be related to the researchers not being able to redo the measurements in the visual QC data set as opposed to their own field measurements. Additionally, the Siikaneva data set, from which the weighting factors for the different measurement classes included in the visual QC exercise were derived, might have contained more measurements that deviate from the expected continuous linear increase in CH_4 concentrations in the chamber than the data sets usually collected by the survey participants. As discussed earlier, this might be related both to the specific measurement setup used but also to the site-specific environmental conditions at Siikaneva bog.

4.4 Implications: Effects of measurement variability and researcher variability and bias on methane fluxes estimated using the chamber-based methods

In this study we performed an expert survey to gain an overview of the approaches used by different researchers to measure, process, and quality control CH_4 fluxes using flux chambers. We analyzed the survey responses to identify the main discrepancies between the approaches used within the flux community as they might cause significant uncertainty when comparing or combining data sets collected and processed by different researchers. Synthesis datasets of chamber CH_4 fluxes

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

have been increasingly used to estimate CH_4 emissions for northern high latitude methane budgets (Kuhn et al., 2021; Treat et al., 2018). Generally, these synthesis datasets show consistent differences among different wetland classes, but additionally, high variability within each wetland class that is attributed to high spatial and temporal variability within the flux measurements that can be partly compensated for by using longer integration times (Treat et al., 2018) or by capturing spatial variability due to microtopography effects (Virkkala et al., 2024).

The results from this expert survey show that differences in methodology may be an additional factor contributing to high variability in methane fluxes found in synthesis datasets. For example, the handling of low positive and negative fluxes can significantly affect estimates of methane budgets particularly for high latitude and upland regions where low CH_4 emissions and/or uptake of CH_4 can be expected during large parts of the year. Discarding low fluxes or setting them to zero can therefore lead to a bias towards higher CH_4 emissions and potentially make the difference between a net annual uptake or a net emission of CH_4 in low flux regions.

Still, the uncertainty caused by different researchers processing a representative data set as derived from our visual QC exercise seems small compared to estimates of the natural spatial and temporal variability in CH_4 fluxes. We estimated an overall variation between flux estimates caused by different researchers choosing different time periods of the same measurement for flux calculation of 17% and a variation in the percentages of measurements passing the visual QC of 28%. These estimates are similar to the mean natural temporal variability of 19% but lower than the mean natural spatial variability in CH_4 fluxes of 88% calculated from autochamber measurements in five temperate and Arctic peatlands by Pirk et al. (2016). Pirk et al. (2016) similarly found that this natural spatial and temporal variability in CH_4 fluxes exceeds the difference between the fluxes estimated using different fit functions. However, it has to be noted that the uncertainty estimates derived in our study consider only the effect of differences in visual QC and do not account for different measurement setups and routines or different fit functions used for flux calculation. Still, this discussion points towards the questions of where and when we introduce the largest error into our flux estimates—is it when we choose our measurement setup and processing approaches or do the location and the timing together with the spatial and temporal resolution of the measurements matter more? Answering this question will help to identify the most relevant starting points for improving the accuracy of flux estimates and help lower uncertainties for flux syntheses. In any case, the survey shows that our human decision-making introduces uncertainties that can obscure natural spatial and temporal variability in CH_4 fluxes. This might make it harder to identify relevant spatial and temporal environmental drivers of CH_4 fluxes, which is crucial for processed based modelling and model development.

5 Conclusions

With this expert survey, we aimed to get an overview of the measurement, data processing and QC techniques that are used within the flux community to estimate CH_4 fluxes from chamber measurements. By teasing out the main differences between the approaches used, we identified major sources of uncertainty in syntheses of data sets processed by different researchers.

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

From our findings we derived starting points for future research to address this uncertainty through refining and substantiating existing recommendations on the measurement setup as well as the data processing and QC approaches.

From the survey responses, most but not all researchers are using a measurement procedure and all of the chamber equipment recommended to avoid chamber artefacts as much as possible. While well founded guidelines on the measurement setup exist in the literature, recommendations on flux calculation and QC approaches are less unambiguous. As a major source of potential bias in flux estimates, we identified widespread uncertainty among the researchers in the handling of chamber measurements showing a pattern in CH₄ concentrations inside the chamber that deviates from the expected linear increase. The reasons presumed for the observed patterns in CH₄ concentrations were manifold which resulted in differing approaches on how to filter a data set as well as on choosing a time period for flux calculation within a measurement, causing uncertainties of 28 and 17%, respectively, when processing a representative data set. The choice of the time period strongly affected the resulting flux estimate especially for nonlinear changes in CH₄ concentrations. Another decision that significantly influenced the magnitude of flux values was whether to include ebullitive CH₄ emission in the flux estimates. Specific questions resulting from the expert survey are: (1) Which part of a nonlinear change in CH₄ concentrations best represents the actual CH₄ flux?; (2) Should ebullitive emissions be included in the CH₄ fluxes estimates from chamber measurements?; (3) Can a linear increase following an ebullition event still be used to calculate the diffusive flux?; (4) How should we deal with small changes in CH₄ concentrations over the course of a chamber measurement?; (5) Under which conditions can net uptake of CH₄ be expected? Besides highlighting the crucial role of complete and details methods descriptions, all of these questions indicate that we need to better understand both the biogeochemical processes underlying the CH₄ fluxes as well as potential chamber artefacts.

Identifying the environmental conditions under which each of the processes can significantly affect the CH₄ concentrations inside the chamber as well as the environmental variables that need to be recorded alongside the gas concentrations in order to characterize these conditions will enable us to make more informed decisions on how to process chamber measurement in the future. Continuing to work towards a standardized measurement setup and measuring procedure for chamber measurements of CH₄ fluxes according to existing and well founded recommendations will reduce the uncertainty between measurements collected by different researches. For data processing and QC, however, rather than being standardized, routines should account for the environmental conditions specific to each research site in order to avoid a bias in individual data sets. Establishing flux community networks for chamber measurements similar to the ones that already exist for the eddy covariance technique will foster a transparent exchange between researchers on measurement and data processing techniques. Such chamber flux networks could build on existing research infrastructures such as the LTER sites, the ICOS sites in Europe and NEON sites in North America. These measures will allow us to reduce the uncertainty in syntheses of data sets processed by different researchers which are urgently needed to estimate CH₄ emissions on large spatial scales. Making individual data sets more comparable and combinable forms a basis for addressing the need of data exchange that was recently acknowledged by the WMO as part of the “Global Greenhouse Gas Watch” (G3W) (WMO, 2023).

Field Code Changed

Formatted: English (United States)

Formatted: English (United States)

1900 *Data availability.* The results of the expert survey described in this paper are available from PANGAEA:
<https://doi.org/10.1594/PANGAEA.971695> (Jentzsch et al., 2024b).

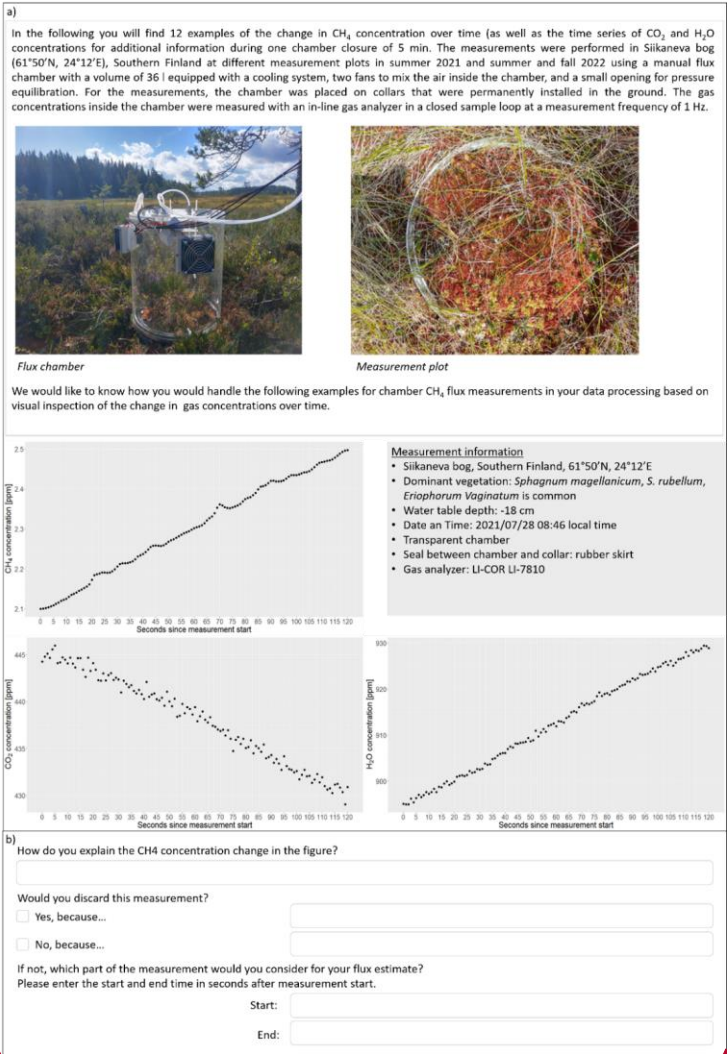
Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Appendix



Formatted: English (United States)

Formatted: Keep with next

Formatted: English (United States)

905 Figure A1: Example of the information provided (a) and the questions asked (b) for the visual QC of one of the 12 chamber measurements under discussion.

Formatted: Keep lines together

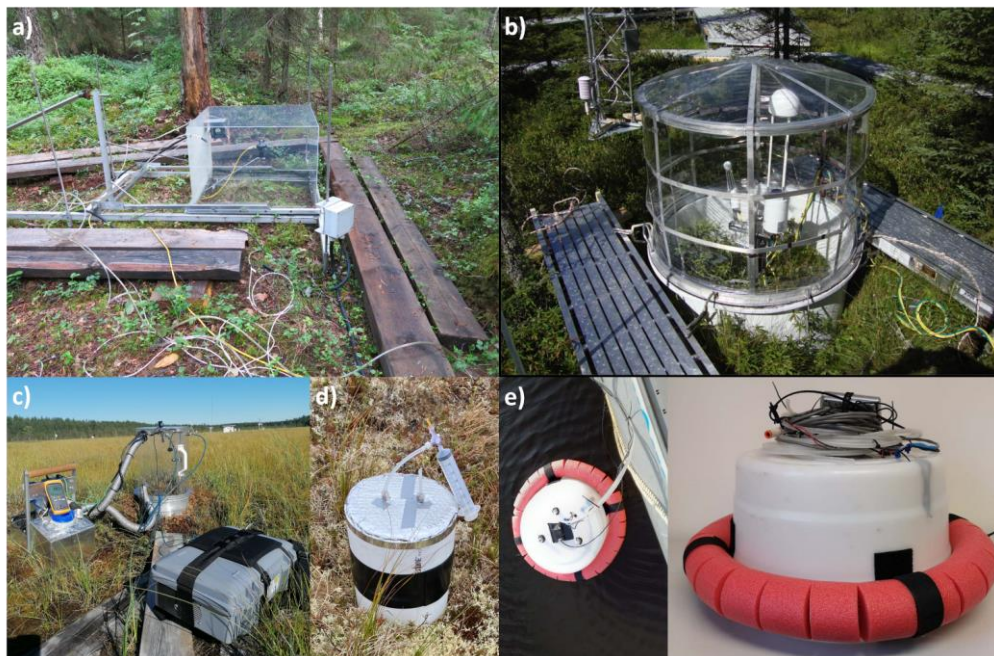


Figure A2: Examples for different chamber setups in different environments. Automated chamber in a boreal forest (a), large manual chamber with gas analyser inside the chamber (b), transparent manual chamber with in-line gas analyser and cooling unit in a boreal fen (c), opaque manual chamber with a syringe for manual gas sampling and a tube for pressure equilibration (d), floating chamber connected to in-line gas analyser and deployed from a boat for aquatic measurements (e).

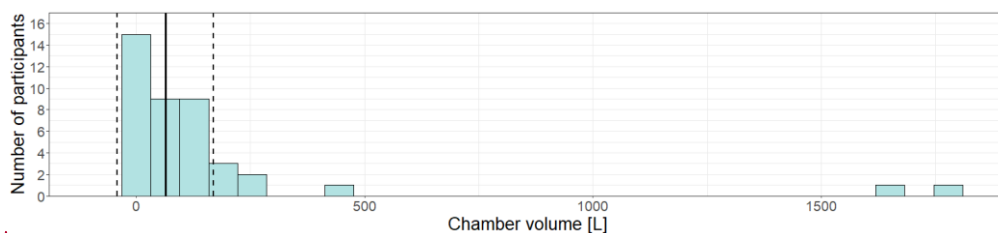
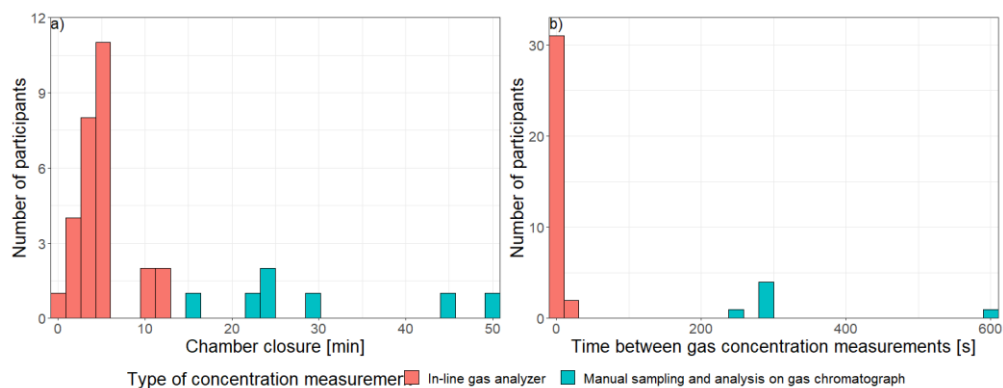
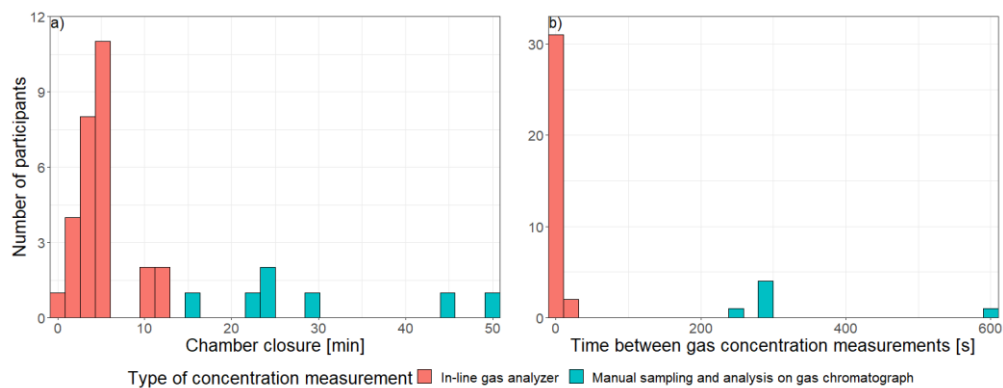


Figure A3: Histogram of the chamber volumes. Black solid line: median, black dashed lines: IQR.



Formatted: English (United States)

Formatted: Keep with next

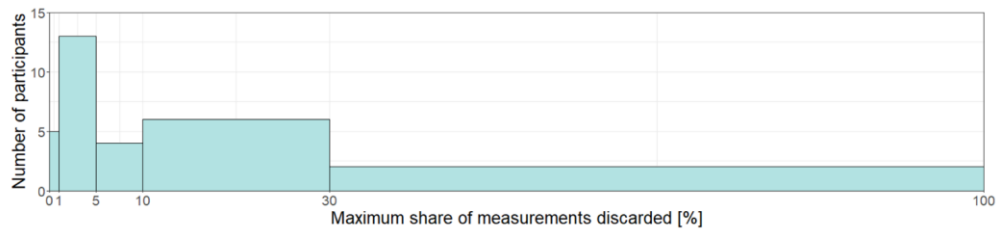


Formatted: English (United States)

Figure A4: Duration of chamber closure (a) and frequency at which the gas concentration inside the chamber is recorded, as time between two measurements (b) by type of concentrations measurement.

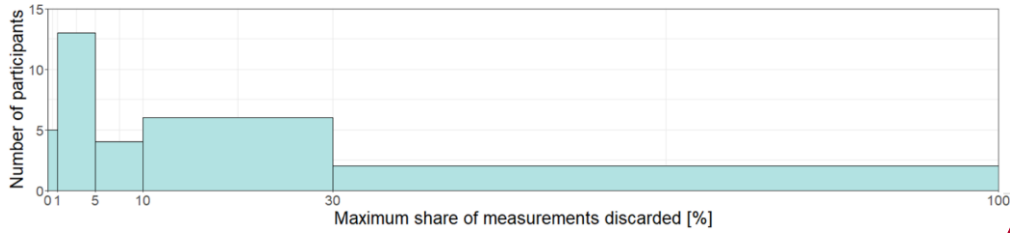
Formatted: Keep lines together

Formatted: English (United States)



Formatted: English (United States)

Formatted: Keep with next



Formatted: English (United States)

Figure A5: Maximum percentage of measurements that the survey participants usually discard from their own data sets.

1925

1930

1935

1940

Table A1: Number and percentage of occurrence of identified measurement scenarios in a data set of chamber measurements collected at Siikaneva bog, Southern Finland, in 2021 and 2022. Categorization of the 12 measurement examples used in the visual QC exercise into these measurement scenarios and weighting factors calculated for each measurement example for the extrapolation of CVs derived from the 12 measurement examples to a representative data set (Table A2). Weighting factors were derived as the relative occurrence of the respective measurement scenario in the Siikaneva data set (788 measurement in total) divided by the number of occurrences in the visual QC exercise.

Measurement scenario	Linear increase		Nonlinear increase – decreasing slope			Initial jump	Jump(s)			Nonlinear increase – increasing slope	Inconsistent trend	Linear decrease	Low variation
Occurrence in Siikaneva data set (%)	468 (59.4)		144 (18.3)			66 (8.4)	62 (7.9)			25 (3.2)	16 (2.0)	4 (0.5)	3 (0.4)
Measurement ID	VOC1	VOC2	VOC4	VOC5	VOC9	VOC7	VOC8	VOC12	VOC10	VOC11	VOC3	VOC6	
Weight (%)	29.7	29.7	6.1	6.1	6.1	8.4	4.0	4.0	3.2	2.0	0.5	0.4	

Table A2: Visual QC exercise (blue shading): Fluxes in mg CH₄ m⁻² d⁻¹ calculated for each example chamber measurement (columns) by each survey participant (rows) based on the time periods chosen for flux calculation in the visual QC part of the survey, by each survey participant. Coefficients of variance (CV) are given for each example measurement (CV_{ex}) and each measurement scenario (CV_{scenario}) across the participants. “NR” indicates that a participant did not respond to the respective measurement example, so neither kept nor discarded the measurement and “D” indicates that a participant discarded a measurement example. Percentage of measurements kept for flux calculation (purple shading): Percentage of measurement examples within the visual QC exercise kept for flux calculation by participant (Kept_{QC}, relative to total number of measurement examples that a participant responded to), measurement example (Kept_{ex}), and measurement scenario (Kept_{scenario}). Extrapolation to Siikaneva data set and uncertainty estimates (red shading): Extrapolation of the visual QC results to the entire Siikaneva data set through weighting by the frequency of occurrence of each measurement scenario in the data set (Table A1X). Answers from participants who gave no response to more than one measurement example were excluded from the calculations of Kept_{QC} and Kept_{Siikaneva}. Participants who did not participate at all in the visual QC exercise have been excluded from the table.

Measurement scenario	Linear increase		Nonlinear increase – decreasing slope			Initial jump	Jump(s)			Nonlinear increase – increasing slope	Inconsistent trend	Linear decrease	Low variation		
Measurement ID	VOC1	VOC2	VOC4	VOC5	VOC9	VOC7	VOC8	VOC12	VOC10	VOC11	VOC3	VOC6	Kept _{QC} (%)	Kept _{Siikaneva} (%)	
1	67.31	3204.96	68.56	171.47	1102.58	351.70	2183.91	87.58	137.75	127.25	-8.99	14.17	100	100	
2	66.80	3169.76	D	D	1049.99	377.13	D	D	131.51	D	D	D	42	77	
3	68.68	3367.11	73.95	169.61	1036.42	371.20	D	88.95	158.75	-18.64	-10.78	13.96	92	96	
4	67.02	3294.94	66.92	171.47	1790.25	350.85	3117.04	482.53	127.40	D	-11.16	D	83	98	
5	67.02 ^a	3166.90 ^a	32.80 ^a	67.90 ^a	1751.88 ^a	D	D	D	116.73 ^a	D	-11.16 ^a	0	67	81	
6	66.34	3156.77	65.24 ^{a,b}	61.26	1535.05	D	D	275.62	D	D	D	D	50	52	
7	67.17	3067.51	61.86	D	1057.30	D	2031.93	D	133.38	D	D	D	50	79	
8	67.02	3169.76	D	D	1091.37	378.93	D	D	116.73	D	-11.16	D	50	78	
9	67.02	3169.76 ^a	71.13	67.90 ^{a,b}	1751.88 ^a	D	D	D	D	D	-11.08	0	58	79	

Formatted: Keep with next, Keep lines together

Formatted Table

Formatted: English (United States)

Formatted: English (United States)

Formatted

Formatted: English (United States)

Formatted: English (United States)

Formatted: Keep with next, Keep lines together

Formatted

Formatted: English (United States)

Formatted

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted

Formatted: Right

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

1960 participants additionally noticed slight deviations from the linear behavior of the CH₄ concentrations. Minor jumps in the CH₄ concentration in VQC1 were mentioned by 17 participants (61%), which they related to CH₄ ebullition (9 participants), insufficient mixing due a defective fan (3 participants), wind (1 participant), wind-induced pressure changes (1 participant), changes in atmospheric pressure influencing the ground diffusion rates and/or atmospheric pressure gradient (1 participant), boundary layer disturbance (1 participant), leakage (2 participants), disturbance (1 participant) caused by chamber placement or footsteps (1 participant). For VQC2, half of the participants pointed out a decrease in the slope of CH₄ concentrations starting between 250 and 260 s after the chamber closure, 21% of whom also noticed a simultaneous decrease in the slope of CO₂ concentrations. As explanations the participants mentioned saturation of the chamber headspace decreasing the concentration gradient over time (5 participants), a build-up of pressure (2 participants) potentially due to a defective pressure valve towards the end of the measurement (1 participant), a change in temperature over the course of the measurement (1 participant), or a small leak (1 participant) probably combined with windy conditions (1 participant). Many participants

1970 furthermore discussed the change in CO₂ and H₂O concentrations over the time of the chamber closure. For VQC1, three participants mentioned that the CO₂ and H₂O concentrations show a linear change, two of whom concluded that there was no air leaking from the chamber. Three participants on the other hand were concerned about the H₂O measurements due to the high and increasing concentrations, and due to an assumed saturation and therefore decreasing slope towards the end of the measurement. Leakage from the chamber was suspected by three participants, two of whom explained this presumption with

1975 vegetation overgrowing the collar and one with the use of a less airtight rubber seal as opposed to a water seal. For VQC2, 18% of the participants picked up on the drop in H₂O concentrations occurring around 40 s after the chamber closure, 40% of whom additionally mentioned a simultaneous change in the slope of CO₂ concentrations. Their reasoning included water condensing on the chamber walls and changing light conditions. Few participants decided to discard the two measurement examples. Measurement VQC1 was discarded by 4 participants (14%) suspecting CH₄ ebullition or stating that the starting concentrations of CO₂ were too high above ambient concentrations or that all chamber measurements generally need to be

1980 shaded. One participant excluded VQC2 due to an assumed saturation effect and one additional participant mentioned ebullition and a high initial concentration of CH₄ as potential reasons to exclude the measurement from flux calculations. 86% and 89% of the participants decided to keep VQC1 and VQC2 for flux calculation, respectively, due to the consistent linear increase in CH₄ concentrations without clear indications of significant disturbances or any malfunctioning of the instruments.

1985 For VQC1, the participants further supported their decision with the linear change in CO₂ and H₂O concentrations making leakage from the chamber unlikely as well as with near-ambient CH₄ concentrations at the measurement start. For both VQC1 and VQC2 most participants who gave start and end times for flux calculation chose the middle part of the measurement, discarding the beginning and the end without mentioning a specific reason. The remaining participants considered the CO₂ and/or H₂O concentrations in their choice of the time period for curve fitting. For VQC1, three participants chose the beginning

1990 of the measurement only, resulting in slightly higher flux estimates, two of whom assumed that H₂O saturation diminished the increase in CH₄ concentrations towards the end of the measurement. For VQC2, some participants acknowledged the strong drop in H₂O concentrations. Having no further information on potential reasons three of them decided not to let this unexpected

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

1995 behavior in H₂O concentrations make them discard the CH₄ measurements while other participants reacted by excluding the
time of the drop in H₂O concentrations from their calculation of the CH₄ flux through either using the part of the measurement
after the drop (7 participants) or before the drop (1 participant). 61% of the 23 participants who entered start and end times for
flux calculation discarded the end of the measurement where CH₄ and CO₂ concentrations increased at a lower rate, resulting
in slightly higher flux estimates above 3200 mg CH₄ m⁻² d⁻¹. Two participants suggested to use a nonlinear fit which one of
them specified as exponential.

2000

Formatted: English (United States)

Formatted: English (United States)

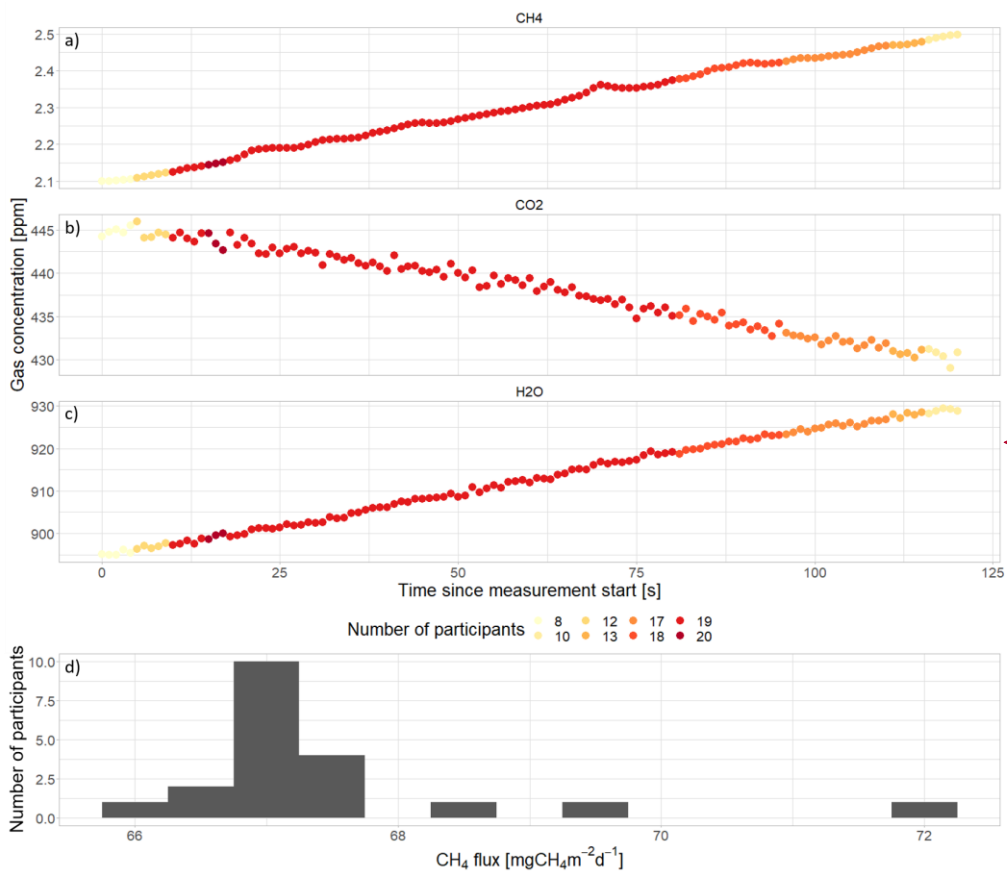


Figure A1.1: Measurement example VQC1 of a small linear increase in CH₄ concentrations over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colors of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

Formatted: English (United States)

Formatted: Keep with next

Formatted: Keep lines together

Formatted: English (United States)

Formatted: English (United States)

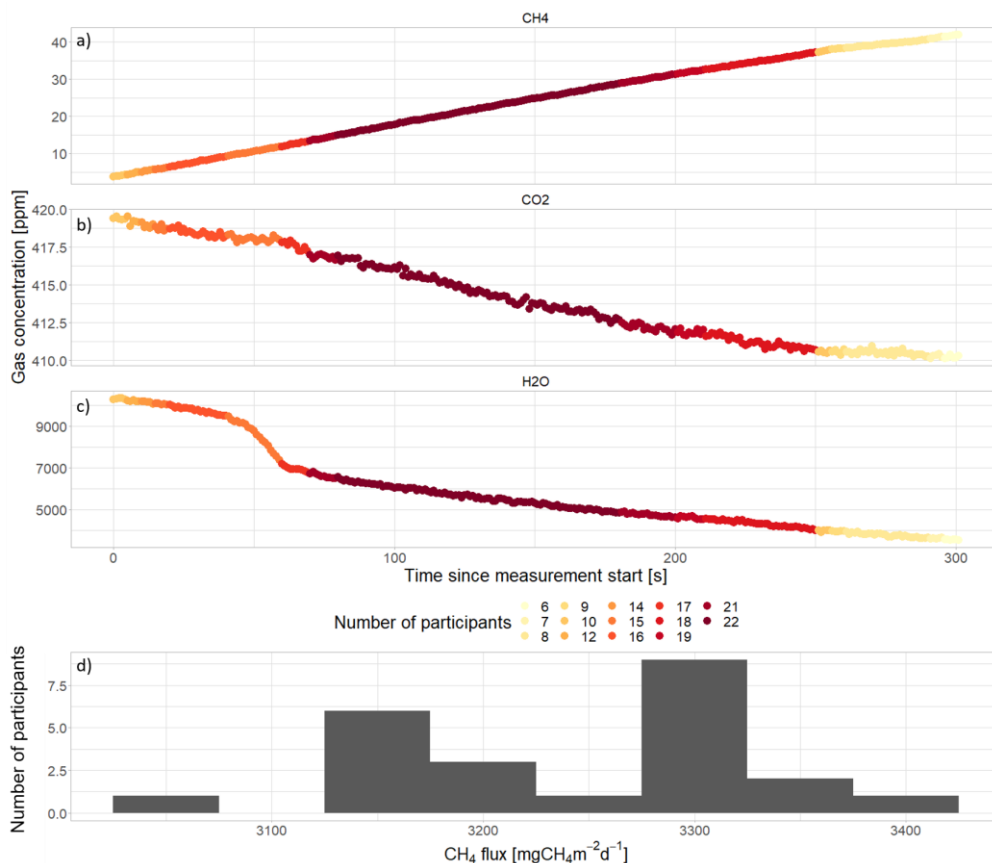


Figure A1.2: Measurement example VOC2 of a strong linear increase in CH₄ concentrations over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colors of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

A1.2 Nonlinear increase – decreasing slope

In the visual QC exercise, we included three examples of measurements that feature a nonlinear increase in CH₄ concentrations during the chamber closure with the rate of increase flattening out over time. Two examples show a small

Formatted: English (United States)

Formatted: Keep with next

Formatted: Keep lines together

Formatted: English (United States)

Formatted: English (United States)

Formatted: Heading 2

Formatted: Indent: First line: 0,5 cm

nonlinear increase in CH₄ concentrations (measurement IDs VQC4 and VQC5, Figures A1.3 and A1.4) simultaneous with linearly decreasing CO₂ concentrations. H₂O concentrations increased over the time of the chamber closure in VQC4 but decreased in VQC5. The third example (measurement ID VQC9, Figure A1.5) shows a stronger increase in CH₄ concentrations with intermittent jumps, linearly increasing CO₂ concentrations and H₂O concentrations that fluctuate without a clear trend.

The participants classified the CH₄ measurements in VQC5 as a small flux that resulted from a balance between CH₄ production and oxidation while VQC9 was identified as large emission indicating a CH₄ hotspot. The majority of the participants (85%, 85%, and 81%) discussed the nonlinear behavior of the CH₄ concentrations in VQC4, VQC5, and VQC9, respectively, offering various explanations for the decreasing rate of increase over time that were mainly related to chamber saturation, chamber leakage or an initial disturbance (Table A1.1).

Most participants (9, 10, and 8) suspected a saturation of the chamber headspace, while two participants stated that saturation was unlikely to be reached during a measurement as short as VQC4 (330 seconds) and one participant explicitly mentioned that the changing slope in VQC9 did not look like a saturation effect. Just as many participants (9) suggested leakage through a weakening seal as the reason for the decreasing slope in VQC4, as supported by the simultaneously decreasing slope in CO₂ and H₂O concentrations while other participants explicitly stated that CO₂ and H₂O concentrations did not indicate a leak in this measurement. Due to the consistently linear CO₂ concentrations in VQC5 and VQC9, only one participant each suspected leakage during these measurement examples. For VQC4, three participants further suspected that the high H₂O concentrations at the end of the measurement influenced the CH₄ measurements, for example through condensation inside the chamber or in the gas flow line, one participant suggested a varying performance of the chamber fan, and two participants assumed that the nonlinearity was a phenomenon specific to *Sphagnum* moss. One participant more generally suggested that vegetation effects changed over the course of the measurement due to changing light conditions, affecting the CO₂ and H₂O concentrations in VQC4 and the CH₄ concentrations in VQC5.

Besides a saturation effect or a weakening seal that would cause a decreasing slope in CH₄ concentrations towards the end of the measurement, many participants (3, 6 and 8) suggested that an initial disturbance such as ebullition triggered by the chamber placement had caused the stronger increase in the beginning of measurement examples VQC4, VQC5, and VQC9. For VQC9, 31% of the participants additionally pointed out minor fluctuations superimposed on the overall nonlinear increase in CH₄ concentrations. Two thirds of them referred to the fluctuations as minor ebullition events while the others suggested episodic leakage from the chamber potentially caused by gusts of wind lifting the chamber sides or a malfunctioning pressure gauge. One participant pointed out that the CH₄ fluctuations cooccurred with fluctuations in the H₂O concentrations and therefore suspected an instrument issue that could be related to spikes in the instrument cavity pressure.

The nonlinearity in the CH₄ concentrations resulted in 15%, 31%, and 12% of the participants deciding to discard the entire measurement example VQC4, VQC5, and VQC9, respectively (Table A1.1). The reasons mentioned for the exclusion of the measurements again reflected the different interpretations of the participants on which part of the measurement represented the real flux. This disagreement shows less strongly in the range of flux estimates since participants who suspected an initial disturbance of the measurement disproportionately often discarded the entire measurement as they assumed that an initial

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

3055

3060

3065

disturbance would also affect the remaining part of the measurement. For VQC4 and VQC5, all 54% of the participants who provided start and end times for flux calculations agreed that the beginning of the measurement should be used for or at least be included in the flux calculation, with three participants suggesting a nonlinear fit for both measurement examples. This resulted in smaller ranges of flux estimates compared to VQC9 (Table A1.1) which instead reflects the fundamentally different interpretations among the participants on which part of the measurement should be used for flux calculation. Here, half of the 21% of participants who gave start and end times for flux calculation chose a later part of the measurement where CH₄ concentrations appeared linear over a longer time period. This resulted in lower flux estimates (between 1000 and 1200 mg CH₄ m⁻² d⁻¹) compared to the flux estimates larger than 3500 mg CH₄ m⁻² d⁻¹ derived for the one quarter of participants who instead chose the beginning of the measurement (Figure A1.5).

Table A1.1: Explanations for the nonlinear increase in CH₄ concentrations, reasons to discard, and reasons and ways to keep measurements showing an increase in CH₄ concentrations as given by the participants. The responses were categorized based on the free text entries for measurement examples VQC4, VQC5, and VQC9. The number of responses given in the respective category are provided in brackets.

<u>Explanations for nonlinearity</u>	<u>Reasons to discard</u>	<u>Reasons to keep</u>
Saturation (23)	Saturation (2)	A nonlinear fit can be used (9)
	Nonlinearity – no steady state reached (3)	
Initial disturbance (16)	Initial disturbance biases flux later on (2)	A (linear) part of the curve can still be used (41)
Bad seal / Leakage from the chamber (8)	Bad seal / Leakage from the chamber (4)	
Unsure (12)	Unclear which part of the measurement represents the real flux (3)	No clear disturbance of the measurement (9)
Changing environmental conditions (1)	Changing environmental conditions (1)	Linear trend in CO ₂ concentrations (5)

Formatted: English (United States)

Formatted: English (United States)

Formatted: Keep with next, Keep lines together

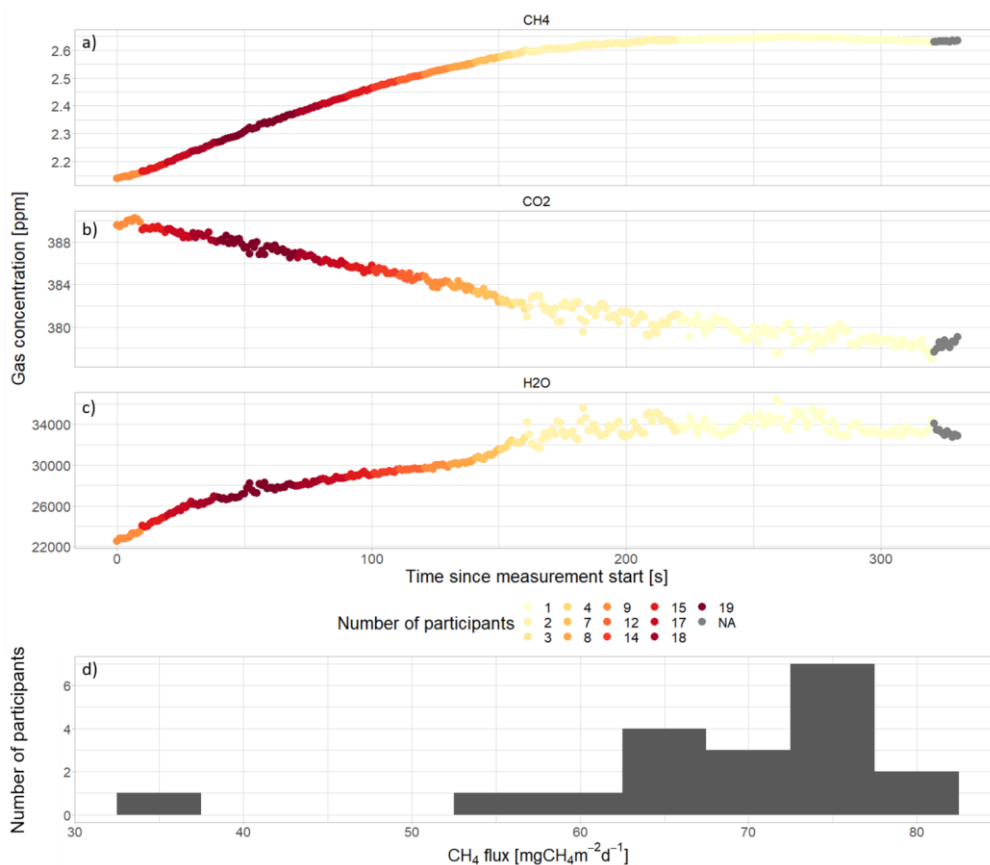


Figure A1.3: Measurement example VOC4 of a small nonlinear increase in CH₄ concentrations with decreasing slope over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colors of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

Formatted: English (United States)

Formatted: Keep with next

Formatted: Keep lines together

Formatted: English (United States)

Formatted: English (United States)

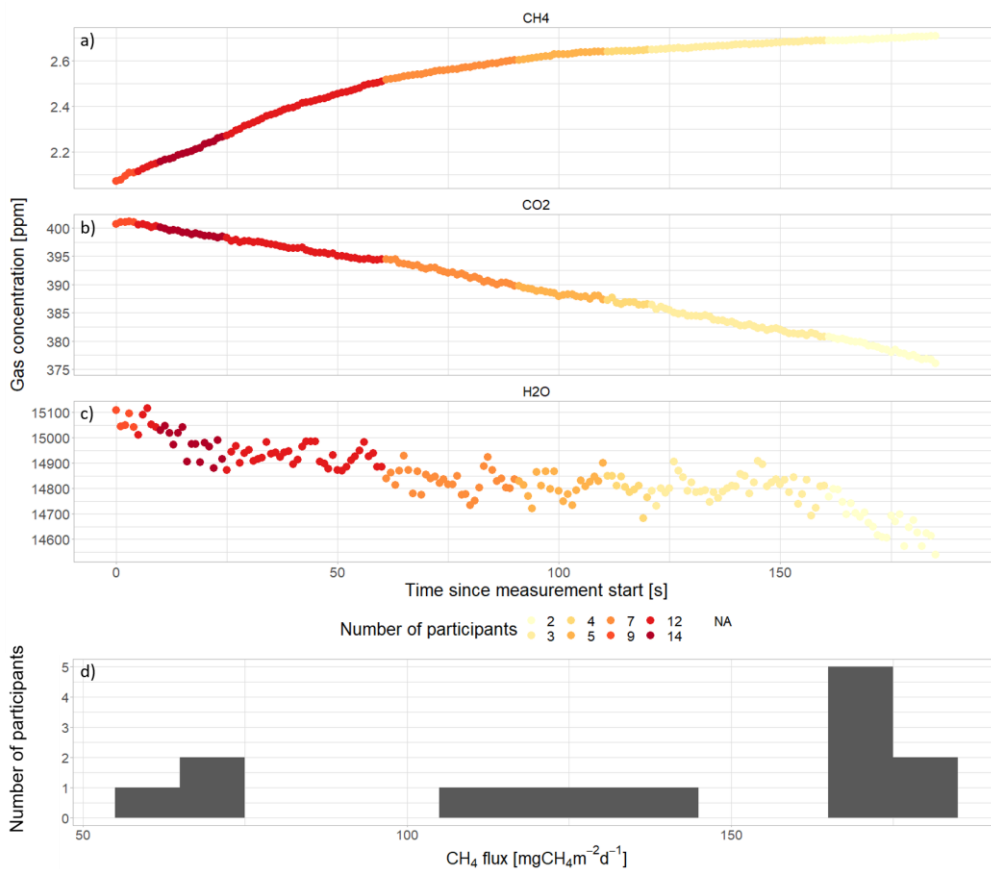


Figure A1.4: Measurement example VQC5 of a small nonlinear increase in CH₄ concentrations with decreasing slope over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colors of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

Formatted: English (United States)

Formatted: Keep with next

Formatted: Keep lines together

Formatted: English (United States)

Formatted: English (United States)

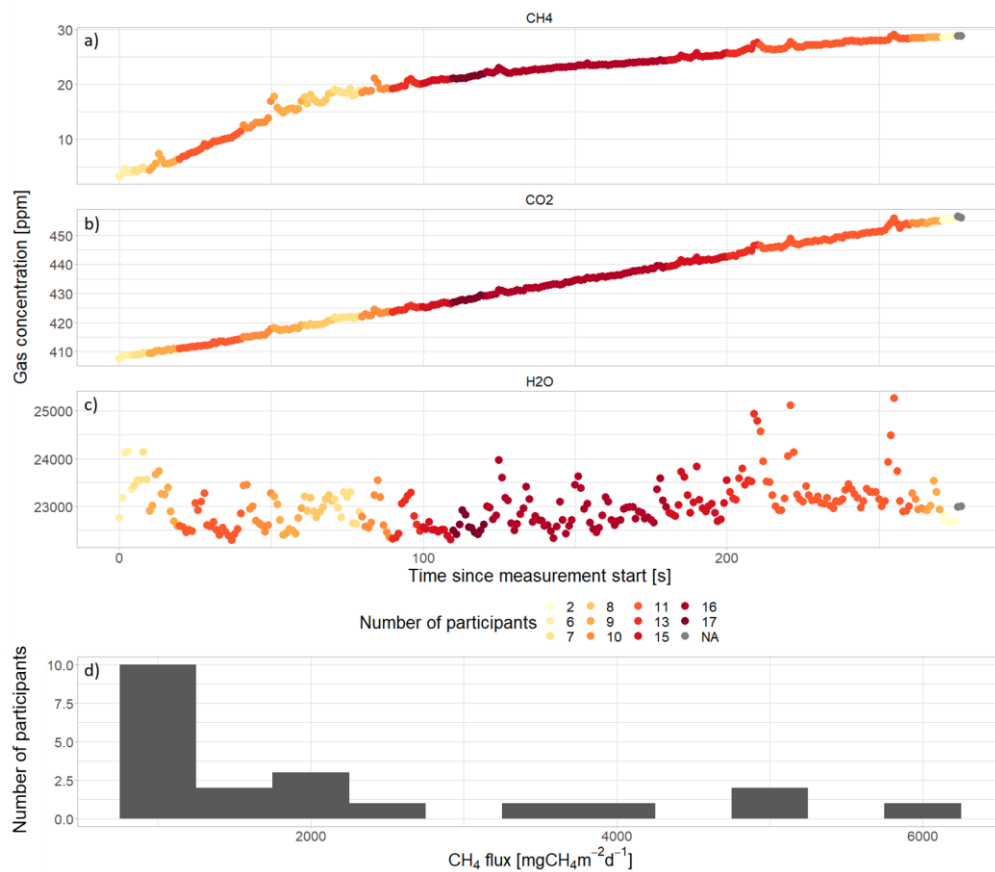


Figure A1.5: Measurement example VQC9 of a strong nonlinear and jumpy increase in CH₄ concentrations with decreasing slope over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colors of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

Formatted: English (United States)

Formatted: Keep with next

Formatted: Keep lines together

Formatted: English (United States)

Formatted: English (United States)

A1.3 Jump(s) at the beginning or in the middle of measurements

In our visual QC exercise, we included three example measurements that showed a relatively linear increase in CH₄ concentrations that was interrupted by one (VQC8, Figure A1.6 and VQC7, Figure A1.7) or more (VQC12, Figure A1.8) sudden increases in the concentration. In one, this occurred at the beginning (VQC7) and in the others, the middle of the measurement (VQC8, VQC12). In examples VQC8 and VQC7, these sudden jumps appeared in all three gases with CH₄ and H₂O concentrations showing a sudden increase while CO₂ concentrations dropped simultaneously. In VQC12, on the contrary, CO₂ and H₂O showed no equivalent to the jumps in the CH₄ concentration. In VQC8, a strong decrease in CH₄ concentrations directly followed the sudden increase, while in VQC7 and VQC12 the concentrations continued to increase at a lower rate starting close to the high concentration level after the jump.

Nearly all (100%, 65%, and 92%) of the participants mentioned the jump(s) in CH₄ concentration when discussing the measurement examples VQC8, VQC7, and VQC12, respectively. For all measurement examples, the majority of these participants explained their observation with episodic events of ebullitive CH₄ emission (VQC8: 65%, VQC7: 88%, VQC12: 92%) with only one participant each suggesting a malfunctioning of the gas analyzer as a reason for the sudden increase in CH₄ concentrations in VQC7 and VQC12. Some (35%) of the participants assuming ebullition stated that the ebullition event in VQC8 was caused by a disturbance and all agreed that the chamber placement caused the ebullition for VQC7. Only one participant (5%) mentioned anthropogenic disturbance as the reason for the ebullition events in VQC12. For VQC8, 12% of the participants pointed out the sudden changes in CO₂ and H₂O concentrations along with the jumps in CH₄. Reasons mentioned by one participant each were a malfunctioning of the gas analyzer and an overpressure caused by the bubble release while another participant suggested the release of gas bubbles with high CH₄ but low CO₂ concentrations as a natural cause for this observation. Similarly, one of the two participants who mentioned the absence of a simultaneous change in the other gases in VQC12, assumed a release of bubbles with high CH₄ concentration but CO₂ concentrations close to ambient conditions due to the different production depths of the two gases. For VQC8, 41% of the participants discussed the decrease in CH₄ concentrations following the assumed ebullition event and suggested leakage of air from the chamber, potentially combined with wind as a potential cause. In the discussion of VQC7, two participants disagreed on the effect of the water table on CH₄ ebullition, one mentioning that in the measurement CH₄ ebullition was more likely to happen because of the high water table while the other stated that ebullition happened despite the high water table, indicating a fundamentally different understanding of the causes of CH₄ ebullition among the participants. Two participants of VQC7 furthermore classified the measurement as an example of strong CH₄ emission which they explained by strong anaerobic CH₄ production related to the high water table and by the vegetation providing substrate for acetoclastic CH₄ production, respectively.

Of the three measurements with jumps in CH₄ concentrations that we included in the visual QC exercise, VQC8 raised the most concern with the highest number of participants excluding the example (Table A2) and with the largest variety of reasons mentioned for the discard, including the inconsistent trend in CH₄ and CO₂ concentrations making them wonder which part of the measurement to use for flux calculation, ebullition affecting the pressure inside the chamber, too much variation in CH₄

Formatted: Heading 2

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

2120 and CO₂ concentrations even after the jump, chamber leakage and too high initial CH₄ and CO₂ concentrations. Leakage was
also suggested by one participant for VQC7, who suspected that Sphagnum moss might have obstructed the chamber seal with
the collar. VQC12 was classified as too short of a measurement by one participant and discarded by another for too high initial
CO₂ concentrations.

2125 There was disagreement among the participants on whether the remaining part of a measurement after a jump in the CH₄
concentration could still be used for flux calculation. For 10 of 11 participants discarding VQC7, the main concern was high
concentrations having a lasting effect on the concentration gradient and thus on the diffusive CH₄ flux during the rest of the
measurement while only one of 21 and 15 participants discarded measurements VQC8 and VQC12 for that reason. For VQC12,
four of the 11 participants who kept the measurement, all of whom also gave start and end times, avoided this problem by
using the beginning of the measurement before the first jump for flux calculation. On the contrary, for VQC8 and VQC7 five
2130 and 14 of the seven and 15 participants who kept the measurement and/or gave start and end times for flux calculation decided
that the measurement after the jump in CH₄ concentrations could still be used for flux calculation, respectively, and five
participants in VQC12 preferred to use the part between the first two jumps because it showed a longer linear increase.

The choice of different time periods for flux calculation resulted in two and three different classes of flux magnitudes for
VQC8 and for VQC7 and VQC12, respectively. The highest flux estimates of more than
2135 3000 mg CH₄ m⁻² d⁻¹, 483 mg CH₄ m⁻² d⁻¹, and 416 mg CH₄ m⁻² d⁻¹ stemmed from the two, one, and two participants who used
the whole measurement example VQC7, VQC8, and VQC12, respectively, for flux calculation because these estimates also
included ebullitive in addition to diffusive CH₄ emissions, reflecting the general disagreement on whether CH₄ ebullition
should directly be included in the flux estimates derived from chamber measurements. For VQC7 and VQC12, the flux
estimates from the participants who excluded the jumps in CH₄ concentration from the time period for flux calculation can
2140 further be split into two classes. For VQC7, nine participants excluded only the very beginning of the measurement, while five
participants only used a later part starting at about 50 s into the measurement when CO₂ concentrations decreased at a higher
rate, resulting in slightly lower CH₄ fluxes. For VQC12, when excluding the jump in CH₄ concentrations the flux estimates
were higher for four participants who chose the measurement period before the first jump, reaching up to 275 mg CH₄ m⁻² d⁻¹
2145 compared to the five participants who chose the longest linear part of the measurement leading to flux as low as 88 mg CH₄ m⁻² d⁻¹. Due to the very linear behavior of the CH₄ concentrations following the initial jump and the higher agreement on the time
period used for flux calculation, the CV of 5% for VQC7 was much lower than for the CVs of 24 and 61% for VQC8 and
VQC12, respectively.

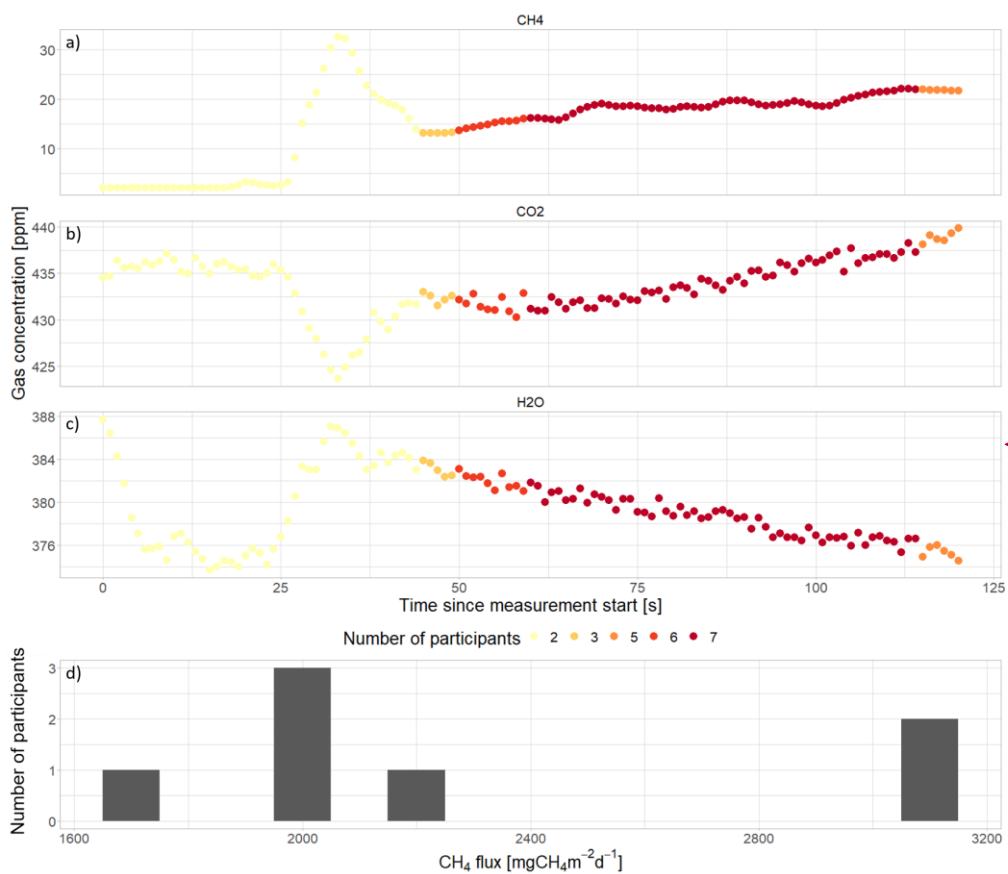


Figure A1.6: Measurement example VOC8 of an overall increase in CH₄ concentrations over the time of the chamber closure after a strong jump in CH₄ concentrations. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colors of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

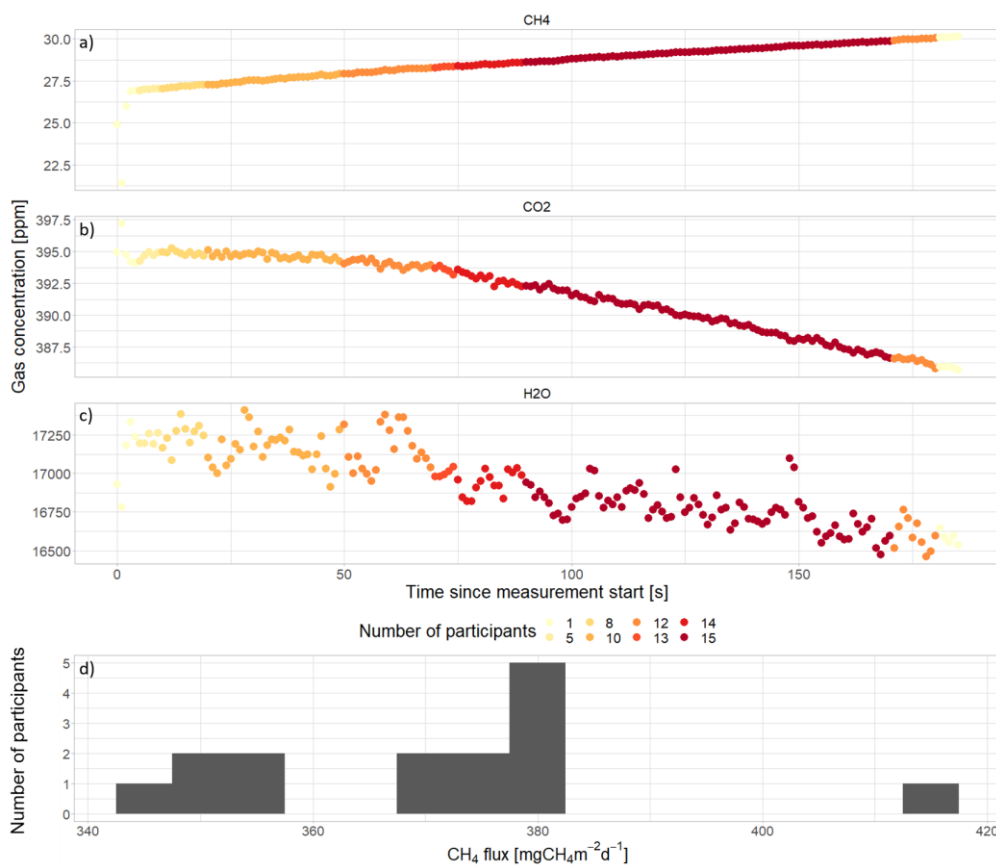


Figure A1.7: Measurement example VOC7 of a linear increase in CH₄ concentrations of the chamber closure after an initial jump in CH₄ concentrations. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colors of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

Formatted: English (United States)

Formatted: Keep with next

Formatted: Keep lines together

Formatted: English (United States)

Formatted: English (United States)



Figure A1.8: Measurement example VOC12 of a linear increase in CH₄ concentrations between repeated jumps in the CH₄ concentration over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colors of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

Formatted: English (United States)

Formatted: Keep with next

Formatted: Keep lines together

Formatted: English (United States)

Formatted: English (United States)

A1.4 Nonlinear increase - increasing slope

One example included in the visual QC exercise showed a nonlinear increase in CH₄ concentrations over the chamber closure with the rate of increase becoming stronger over time (measurement ID VQC10, Figure A1.9). 15% of the participants classified the measurement as a diffusive emission of CH₄ without mentioning further details while 65% discussed the increasing slope in CH₄ concentrations over time, suggesting various reasons that could have caused the observed shape of the curve. The reasons suggested included an initial period of mixing or adjusting, an increase in chamber temperature over time, a disturbance of the measurement plot, a disturbance of the concentration gradient in the soil during chamber placement, an influence of the chamber on plant-mediated CH₄ transport, an incomplete seal of the chamber, incomplete mixing, and an interference with the simultaneously increasing H₂O concentrations. Two participants mentioned that they had not seen such a shape in CH₄ concentrations from chamber measurements before. Regarding the magnitude of CH₄ emissions, three participants pointed out the strong increase in CH₄ concentrations despite the relatively low water table, which they related to plant-mediated CH₄ transport. One participant further mentioned that also the emission of CO₂ was high, indicating warm peat conditions. Two participants mentioned the higher and decreasing CO₂ concentrations in the beginning of the measurement which one of them related to the chamber placement, pushing more gases out of the ground. One participant furthermore mentioned that the chamber seal seemed to be intact.

Six participants decided to discard the measurement, three of whom did so because they could not explain the shape of the curve and stated that the curvature was so strong that the flux estimate would strongly depend on the time period chosen for flux calculation. The three remaining participants mentioned similarly unexpected shapes of CO₂ and H₂O concentrations, higher H₂O concentrations towards the end of the chamber closure which might have interfered with the CH₄ measurements, and high initial CH₄ concentrations as reasons to discard the measurement. 19 participants kept the measurement for flux calculation. The flux estimates for the 17 participants who gave start and end times for flux calculation strongly depended on the time period they chose which in turn depended on their interpretation of the measurement resulting in three distinct classes of flux magnitudes. Two participants decided to use the entire measurement, resulting in intermediate flux estimates of 117 mg CH₄ m⁻² d⁻¹. The majority (13) decided to remove the first 20 to 120 s of the measurement to keep only the more linear part of the CH₄ concentrations in the end, resulting in the highest flux estimates between 125 and 170 mg CH₄ m⁻² d⁻¹. The two remaining participants chose only the linear first 60 or 70 s of the measurement for flux calculation resulting in lower flux estimates of 75 and 76 mg CH₄ m⁻² d⁻¹, respectively, due to the lower rate of increase. Two participants suggested to use a nonlinear fit which one of them specified as exponential.

Formatted: English (United States)
Formatted: Heading 2
Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)
Formatted: English (United States)
Formatted: English (United States)

Formatted: English (United States)
Formatted: English (United States)
Formatted: English (United States)
Formatted: English (United States)

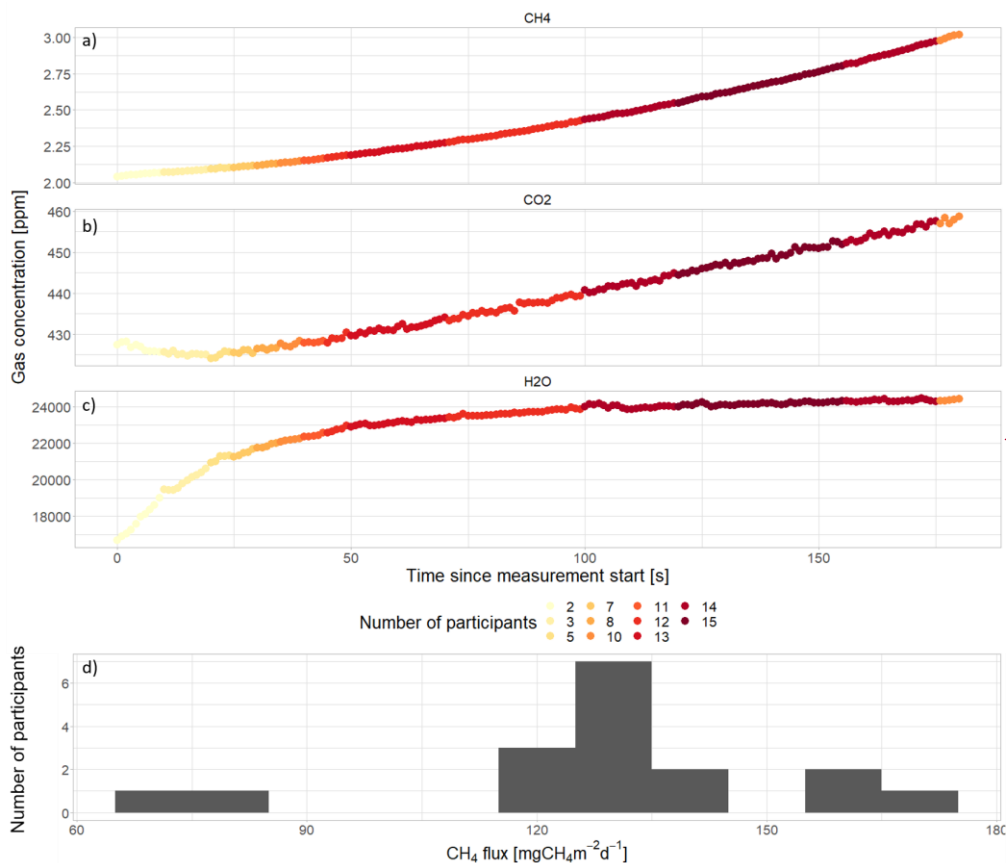


Figure A1.9: Measurement example VOC10 of a small nonlinear increase in CH₄ concentrations with increasing slope over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colors of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

A1.5 Inconsistent trend

One example included in our visual QC exercise showed an inconsistent trend in CH₄ with a change from increasing to decreasing concentrations over the time of the chamber closure (measurement ID VQC11, Figure A1.10). The survey participants disagreed on the reason for this behavior of the CH₄ concentrations. One part of the participants stated that CH₄ oxidation as indicated by the decrease in CH₄ concentrations towards the end of the measurement was unexpected and suggested that measurement issues were responsible for the inconsistent trend in CH₄ concentrations. They had different opinions however on the timing of the disturbance and therefore on which part of the measurement represented the actual CH₄ flux. Some participants suggested an initial disturbance such as CH₄ ebullition caused by the chamber placement while others assumed that the measurement was disturbed at a later point by a problem with the CH₄ analyzer like saturation of the detector or H₂O interference due to the high concentrations towards the end of the measurement and potentially condensation of water vapor, or leakage or a malfunctioning fan after about 50 s into the measurement.

Most participants (66%) discarded the measurement because they missed a consistent trend of sufficient length in the CH₄ concentrations. Since the changing trend was either related to a disturbance or the reason was described as unclear, the participants did not know which part of the measurement to use for the flux calculation. Two participants additionally discarded the measurement because they considered the changes in the CH₄ concentration as too close to zero and another participant mentioned that the CO₂ and H₂O concentrations did not show a steady trend over time either. Some (23%) of the participants decided to keep the measurement for flux calculation, all of whom provided start and end times for flux calculation as well as one additional participant who was uncertain whether to keep or to discard the measurement. The choice of the time periods used for flux calculation depended on the interpretation of the observed pattern in CH₄ concentrations and thus strongly influenced the resulting flux estimate ranging between a CH₄ uptake of -19 mg CH₄ m⁻² d⁻¹ to CH₄ emissions of up to 139 mg CH₄ m⁻² d⁻¹ and splitting the flux histogram into three distinct modes. Two participants chose to keep the entire measurement, resulting in a small positive flux indicating small net CH₄ emission of 8 to 10 mg CH₄ m⁻² d⁻¹. Three participants decided to use the stronger increase in CH₄ concentrations in the beginning of the measurement, resulting in the highest CH₄ emissions between 127 and 139 mg CH₄ m⁻² d⁻¹ while two participants assumed that CH₄ was consumed at the plot, using the later decreasing part of the CH₄ concentrations, resulting in negative flux estimates between -6 and -19 mg CH₄ m⁻² d⁻¹. This resulted in the highest CV among the measurement scenarios, estimated at 138%.

- Formatted: Heading 2
- Formatted: Indent: First line: 0,5 cm
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)
- Formatted: English (United States)

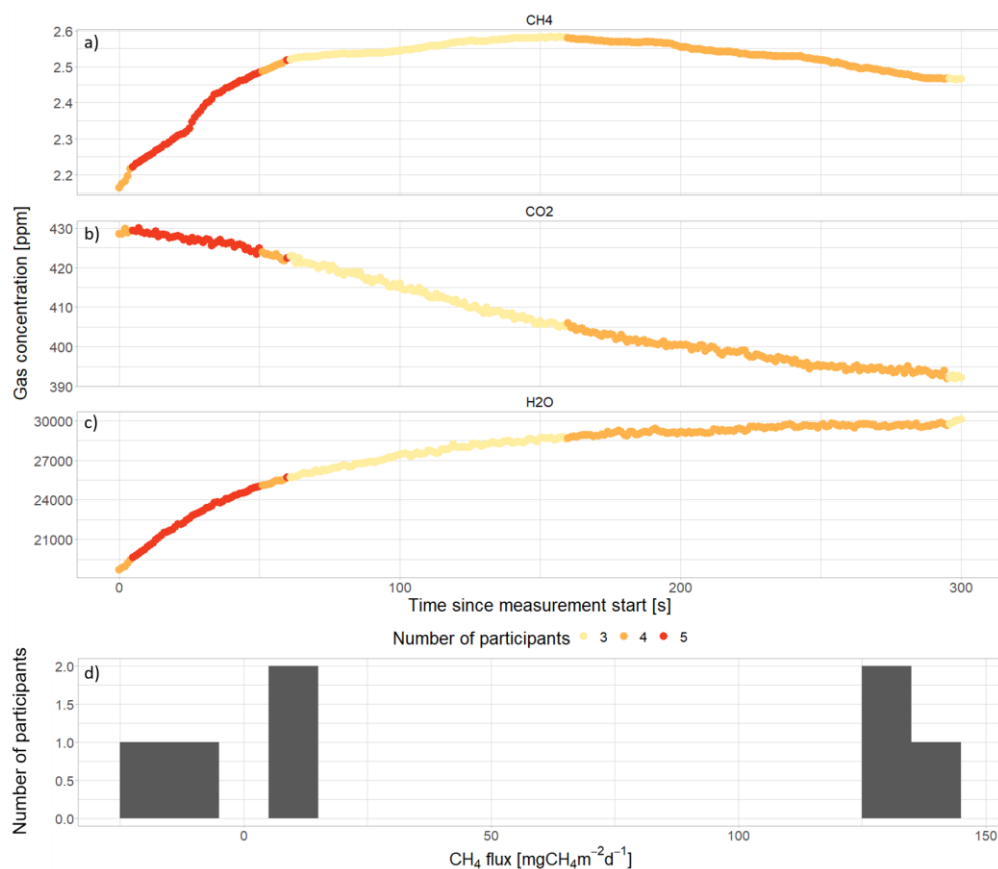


Figure A1.10: Measurement example VQC11 of CH₄ concentrations showing an inconsistent trend over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colors of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

Formatted: English (United States)

Formatted: Keep with next

Formatted: Keep lines together

Formatted: English (United States)

Formatted: English (United States)

A1.6 Linear decrease

One of the measurements in the visual QC exercise showed a small linear decrease in CH₄ concentrations over time (measurement ID VQC3, Figure A1.11). The survey participants largely disagreed on whether this measurement represented a real CH₄ flux. The majority (65%) of the participants assumed real net CH₄ uptake due to CH₄ oxidation dominating over CH₄ production while some (19%) of the participants referred to leakage and too high initial CH₄ concentrations in the chamber as technical problems causing a false apparent uptake of CH₄. The remaining 15% of the participants explicitly stated that they were unsure if the measurement represented a real flux. 23% of the participants more specifically mentioned an inconsistent trend in the CH₄ concentrations referring to three different stages of CH₄ flux or nonlinearities at the beginning and at the end of the measurement. As explanations, they offered initial CH₄ ebullition caused by the chamber placement, changes in the chamber temperature, changes in wind speed combined with chamber leakage, or changes in PAR potentially due to a changing cloud cover or due to condensation inside the chamber indicated by the trend in CO₂ concentrations changing along with the CH₄ trend as well as by high H₂O concentrations.

A slim majority (54%) of the participants discarded the measurement because they did not expect CH₄ uptake in the given environmental (despite the relatively low water table), or because of the inconsistent trend in CH₄ concentrations which makes them unsure which part of the measurement to use for flux calculation, or because of too high initial concentrations of CH₄ and/or CO₂, or because they suspected anthropogenic disturbance from footprints and compacted vegetation or leakage. The flux estimates derived from the start and end times given by 11 of the 12 participants who decided to keep the measurement (46%) differed between the time periods chosen for flux calculation. While five participants chose the entire measurement, resulting in intermediate values of CH₄ uptake, the remaining six participants chose the time period for curve fitting based on the CO₂ concentrations. The middle part of the measurement with linearly decreasing CO₂ concentrations, the beginning of the measurement with stable CO₂ concentrations, and the end of the measurement with linearly increasing CO₂ concentrations were chosen by one, two, and one participant, respectively, while two participants excluded the end of the measurement resulting in strongly negative, lower negative, stronger negative and intermediate CH₄ fluxes, respectively. Overall, the mean of the flux calculated by the 12 experts keeping this flux was 11.36 mg CH₄ m⁻² d⁻¹ with a CV of 17%.

Formatted: Heading 2

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

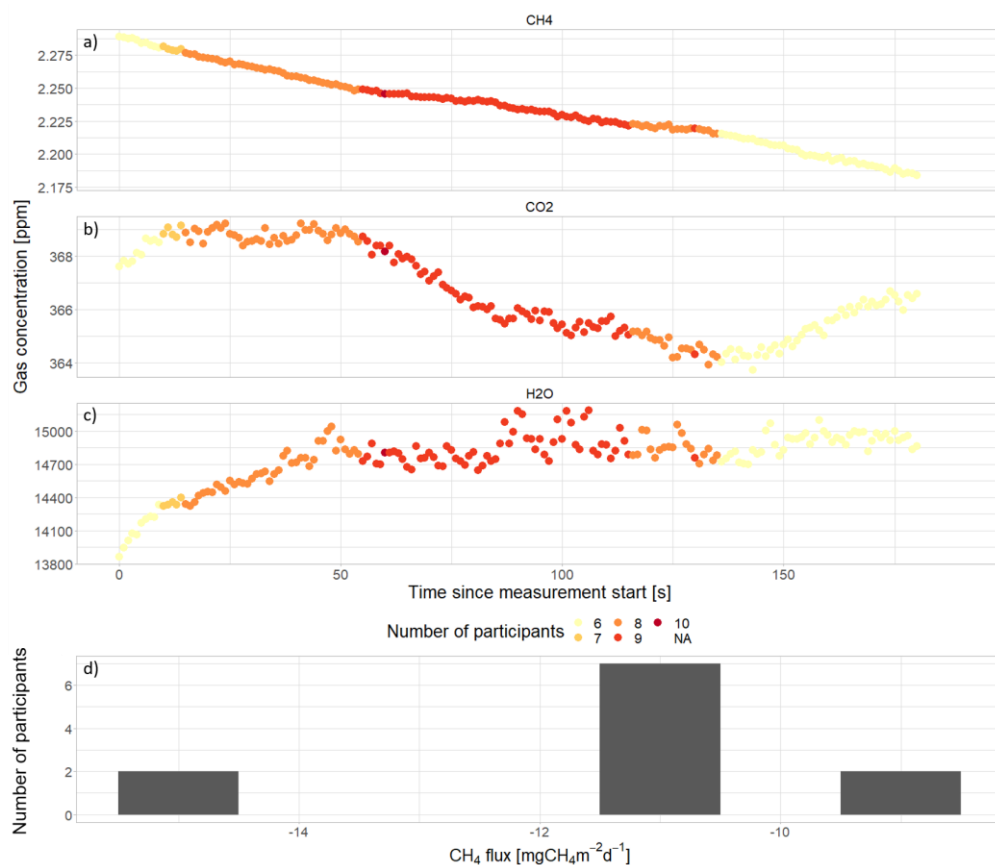


Figure A1.11: Measurement example VOC3 of a small linear decrease in CH₄ concentrations over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colors of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants (d).

Formatted: English (United States)

Formatted: Keep with next

Formatted: Keep lines together

Formatted: English (United States)

Formatted: English (United States)

A1.7 No trend

In the visual QC exercise, we included one measurement example for which the CH₄ concentrations did not show a clear trend and varied only little over the time of the chamber closure (measurement ID VOC6, Figure A1.12). Most participants (69%) noticed the very small change in CH₄ concentrations over the whole measurement but they disagreed on whether the concentration measurements represented a real flux. Half of them suspected a real emission that remained very small because of CH₄ production and oxidation cancelling each other out at a low water table and two more participants called it a “zero flux” where the uncertainty would likely exceed the flux magnitude. Some (39%) of the participants, however, explained the low change in CH₄ concentrations by air leaking from the chamber, two of whom related the leak to vegetation obstructing the chamber seal and one to lateral diffusion into the chamber from the surrounding area.

Some (19%) of the participants furthermore pointed out an inconsistent trend in the CH₄ concentrations which they related to a changing balance between CH₄ production and oxidation over time, noisy measurements due to a low precision of the gas analyzer, or a bad chamber seal combined with wind disturbance. According to one participant the latter was supported by the fluctuations appearing in the concentrations of all three gases, while two other participants mentioned that the CO₂ concentrations looked linear, at least after 30 to 40 s into the measurement, indicating an intact chamber seal.

The majority of the participants (62%) decided to discard the measurement due to leakage from the chamber (38%), a changing trend in the CH₄ concentrations (44%), a too short measurement time (13%), or too high initial concentrations of CH₄ and CO₂ (13%). While two of these participants manually set the CH₄ flux to zero, one participant pointed out that the concentration changes were too large to be below the precision of the instrument so that the measurement should not be accepted as a zero flux. Some (31%) of the participants kept the measurement assuming a small but nonetheless real CH₄ flux and gave start and end times for flux calculation. Half of them discarded the beginning of the measurement as a period of initial equilibration, while the other half kept the entire measurement. The choice of different time periods for flux calculation by the participants resulted in a CV of 57% for this measurement example.

Formatted: Heading 2

Formatted: Indent: First line: 0,5 cm

Formatted: English (United States)

Formatted: English (United States)

Formatted: English (United States)

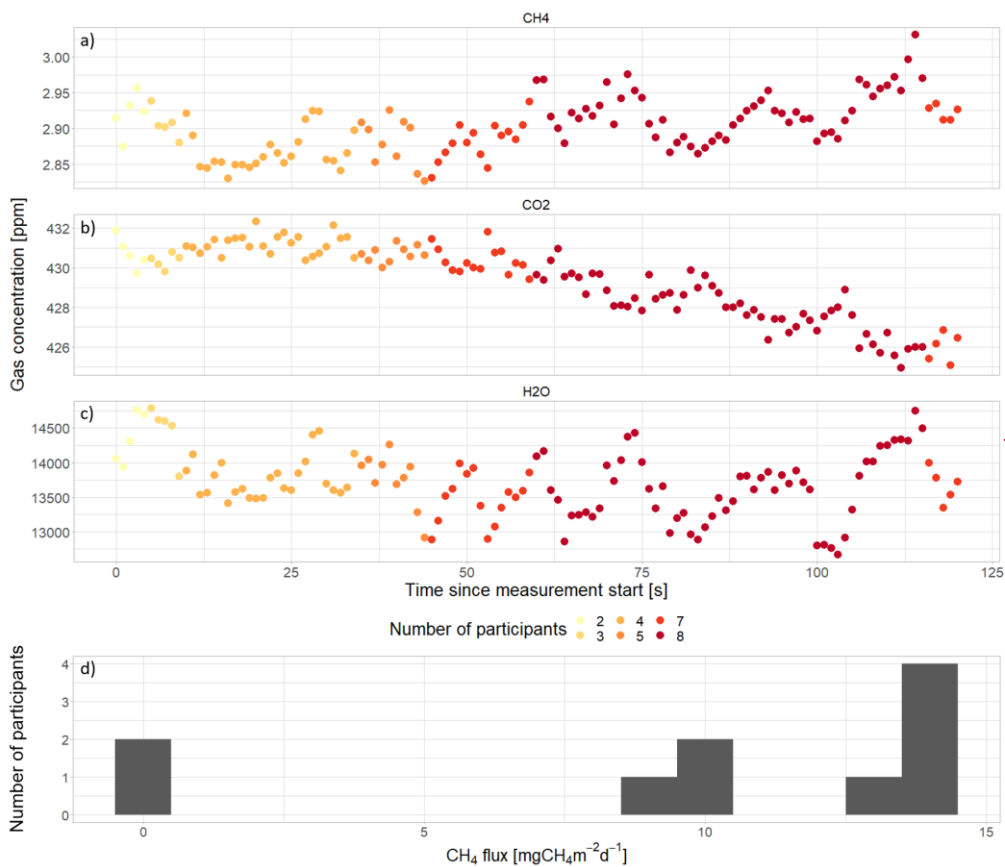


Figure A1.12: Measurement example VOC6 with the CH₄ concentrations showing little variation without a clear trend over the time of the chamber closure. Simultaneous measurements of CH₄ (a), CO₂ (b), and H₂O (c) concentrations over time during chamber closure with the colors of the data points indicating how many participants included the respective data point in the time period that they chose for flux calculation. Only the responses by participants who gave start and end times for flux calculation were considered for this figure. Histogram of CH₄ fluxes calculated based on the time period chosen for flux calculation by the participants including the two participants who set the flux to zero (d).

Formatted: English (United States)

Formatted: Keep with next

Formatted: Keep lines together

Formatted: English (United States)

Formatted: English (United States)

Formatted: Font: Not Bold, English (United States)

Formatted: English (United States)

2305 *Author contributions.* CCT, MF, and KJ conceived the project idea. KJ drafted the questions for the expert survey, set up the final online version of the survey, and collected the Siikaneva data set from which a subset of measurements was used as examples for the visual QC exercise. CCT, LvD, and MF reviewed the survey questions and tested the survey. KJ analysed the survey responses and created the figures. The manuscript was written by KJ and commented on by all authors. CCT supervised the project.

2310 *Competing interests.* The authors declare that they have no conflict of interest.

Acknowledgements. We thank Kathleen Hall, Katey Walter Anthony, Clayton Elder, Vasilii Petrenko for helpful discussions on methods that resulted in the conception of this project. We would like to thank Pamela Baur, Julia Boike, Jill Bubier, Jesper Christiansen, Scott J. Davidson, Lona van Delden, Bo Elberling, Kathleen Hall, Paul Hanson, Nicholas Hasson, Liam Heffernan, Jacqueline Hung, Vytas Huth, Gerald Jurasinski, Sari Juutinen, Masako Kajiura, Evan Kane, Aino Korrensalo, 2315 Elisa Männistö, Nicholas Nickerson, Genevieve Noyce, Frans-Jan Parmentier, Matthias Peichl, Norbert Pirk, Maria Strack, Eeva-Stiina Tuittila, Anna-Maria Virkkala, Carolina Voigt, Lei Wang as well as seven anonymous participants for completing the survey.

Financial support. The contribution of Katharina Jentsch, Lona van Delden, and Claire C. Treat is part of the FluxWIN project, funded with a Starting Grant by the European Research Council (ERC) (ID 851181). The work by Katharina Jentsch was supported by a fellowship of the German Academic Exchange Service (DAAD). CT also acknowledges support from the Pioneer Center Land-CRAFT at AU.

Formatted: Justified

Formatted: English (United States)

Formatted: English (United States)

References

2325 ~~Bain, W.G., Hutyrá, L., Patterson, D.C., Bright, A.V., Daube, B.C., Munger, J.W., and Wofsy, S.C., 2005.~~ Wind-induced error in the measurement of soil respiration using closed dynamic chambers. ~~Agricultural and Forest Meteorology~~ 131, 225–232. ~~https://doi.org/10.1016/j.agrformet.2005.06.004, 2005.~~
~~Bond-Lamberty, B., and Thomson, A., 2010.~~ A global database of soil respiration data. ~~Biogeosciences~~ 7, 1915–1926. ~~https://doi.org/10.5194/bg-7-1915-2010, 2010.~~
~~Bond-Lamberty, B., and Thomson, A., 2010.~~ Temperature-associated increases in the global soil respiration record. ~~Nature~~ 464, 579–582. ~~https://doi.org/10.1038/nature08930, 2010.~~
2330 ~~Christiansen, J.R., Korhonen, J.F.J., Juszczak, R., Giebel, M., and Pihlatie, M., 2011.~~ Assessing the effects of chamber placement, manual sampling and headspace mixing on CH₄ fluxes in a laboratory experiment. ~~Plant Soil~~ 343, 171–185. ~~https://doi.org/10.1007/s11104-010-0701-y, 2011.~~
~~Christiansen, J.R., Outhwaite, J., and Smukler, S.M., 2015.~~ Comparison of CO₂, CH₄ and N₂O soil-atmosphere exchange measured in static chambers with cavity ring-down spectroscopy and gas chromatography. ~~Agricultural and Forest Meteorology~~ 211–212, 48–57. ~~https://doi.org/10.1016/j.agrformet.2015.06.004, 2015.~~
2335 ~~Clough, T.J., Rochette, P., Thomas, S.M., Pihlatie, M., Christiansen, J.R., and Thorman, R.E., 2020.~~ Global Research Alliance N₂O chamber methodology guidelines: Design considerations. ~~J of Env Quality~~ 49, 1081–1091. ~~https://doi.org/10.1002/jeq2.20117, 2020.~~

Formatted: English (United States)

- Conen, F. ~~and~~; Smith, K.A.: ~~1998~~. A re-examination of closed flux chamber methods for the measurement of trace gas emissions from soils to the atmosphere. *European J Soil Science* 49, 701–707. <https://doi.org/10.1046/j.1365-2389.1998.4940701.x>. ~~1998~~.
- de Klein, C.A. ~~and~~; Harvey, M.J., 2012. Nitrous oxide chamber methodology guidelines de Klein, C. A., & Harvey, M. J. Fiedler, J., Fuß, R., Glatzel, S., Hagemann, U., Huth, V., Jordan, S., Jurasinski, G., Kutzbach, L., Maier, M., Schäfer, K., Weber, T., ~~and~~ Weymann, D.: ~~2022~~. BEST PRACTICE GUIDELINE Measurement of carbon dioxide, methane and nitrous oxide fluxes between soil-vegetation-systems and the atmosphere using non-steady state chambers. Arbeitsgruppe Bodengase, Deutsche Bodenkundliche Gesellschaft. <https://doi.org/10.23689/FIDGEO-5422>, ~~2012~~.
- Forbrich, I., Kutzbach, L., Hormann, A., ~~and~~ Wilmking, M.: ~~2010~~. A comparison of linear and exponential regression for estimating diffusive CH₄ fluxes by closed-chambers in peatlands. *Soil Biology and Biochemistry* 42, 507–515. <https://doi.org/10.1016/j.soilbio.2009.12.004>, ~~2010~~.
- Frenzel, P. ~~and~~ Karofeld, E.: ~~2000~~. CH₄ emission from a hollow-ridge complex in a raised bog: The role of CH₄ production and oxidation. *Biogeochemistry* 51, 91–112. <https://doi.org/10.1023/A:1006351118347>, ~~2000~~.
- Healy, R.W., Striegl, R.G., Russell, T.F., Hutchinson, G.L., ~~and~~ Livingston, G.P.: ~~1996~~. Numerical Evaluation of Static-Chamber Measurements of Soil-Atmosphere Gas Exchange: Identification of Physical Processes. *Soil Science Society of America Journal* 60, 740–747. <https://doi.org/10.2136/sssaj1996.03615995006000030009x>, ~~1996~~.
- Hoffmann, M., Schulz-Hanke, M., Garcia Alba, J., Jurisch, N., Hagemann, U., Sachs, T., Sommer, M., ~~and~~ Augustin, J.: ~~2017~~. A simple calculation algorithm to separate high-resolution CH₄ flux measurements into ebullition- and diffusion-derived components. *Atmos. Meas. Tech.* 10, 109–118. <https://doi.org/10.5194/amt-10-109-2017>, ~~2017~~.
- Holland, E.A., Robertson, G.P., Greenberg, J.P., Groffman, P.M., Boone, R.D., ~~and~~ Gosz, J.R.: ~~1999~~. Soil CO₂, N₂O, and CH₄ exchange, in: ~~Robertson, G.P., Coleman, D.C., Bledsoe, C.S., Sollins, P. (Eds.), Standard Soil Methods for Long-Term Ecological Research, edited by: Robertson, G.P., Coleman, D.C., Bledsoe, C.S., and Sollins, P., Oxford University Press, Oxford, UK, pp. 185–201, 1999.~~
- Hutchinson, G.L. ~~and~~ Livingston, G.P.: ~~2001~~. Vents and seals in non-steady-state chambers used for measuring gas exchange between soil and the atmosphere. *European J Soil Science* 52, 675–682. <https://doi.org/10.1046/j.1365-2389.2001.00415.x>, ~~2001~~.
- Jentzsch, K., Männistö, E., Maruschak, M.E., Korrensalo, A., Van Delden, L., Tuittila, E.-S., Knoblauch, C., ~~and~~ Treat, C.C.: ~~2024a~~. Shoulder season controls on methane emissions from a boreal peatland. *Biogeosciences* 21, 3761–3788. <https://doi.org/10.5194/bg-21-3761-2024>, ~~2024a~~.
- Jentzsch, K., Van Delden, L., Fuchs, M., ~~and~~ Treat, C.C.: ~~2024b~~. An expert survey on chamber measurement techniques and approaches for calculation and quality control of methane fluxes. <https://doi.org/10.1594/PANGAEA.971695>, ~~2024b~~.
- Jian, J., Vargas, R., Anderson-Teixeira, K., Stell, E., Herrmann, V., Horn, M., Kholod, N., Manzon, J., Marchesi, R., Paredes, D., ~~and~~ Bond-Lamberty, B.: ~~2020~~. A restructured and updated global soil respiration database (SRDB-V5). <https://doi.org/10.5194/essd-2020-136>, ~~2020~~.
- Kirschke, S., Bousquet, P., Ciais, P., Saunois, M., Canadell, J.G., Dlugokencky, E.J., Bergamaschi, P., Bergmann, D., Blake, D.R., Bruhwiler, L., Cameron-Smith, P., Castaldi, S., Chevallier, F., Feng, L., Fraser, A., Heimann, M., Hodson, E.L., Houweling, S., Josse, B., Fraser, P.J., Krummel, P.B., Lamarque, J.-F., Langenfelds, R.L., Le Quéré, C., Naik, V., O'Doherty, S., Palmer, P.I., Pison, I., Plummer, D., Poulter, B., Prinn, R.G., Rigby, M., Ringeval, B., Santini, M., Schmidt, M., Shindell, D.T., Simpson, I.J., Spahni, R., Steele, L.P., Strode, S.A., Sudo, K., Szopa, S., Van Der Werf, G.R., Voulgarakis, A., Van Weele, M., Weiss, R.F., Williams, J.E., ~~and~~ Zeng, G.: ~~2013~~. Three decades of global methane sources and sinks. *Nature Geosci* 6, 813–823. <https://doi.org/10.1038/ngeo1955>, ~~2013~~.
- Kuhn, M.A., Varner, R.K., Bastviken, D., Crill, P., MacIntyre, S., Turetsky, M., Walter Anthony, K., McGuire, A.D., ~~and~~ Olefeldt, D.: ~~2021~~. BAWLD-CH₄: a comprehensive dataset of methane fluxes from boreal and arctic ecosystems. *Earth Syst. Sci. Data* 13, 5151–5189. <https://doi.org/10.5194/essd-13-5151-2021>, ~~2021~~.
- Kutzbach, L., Schneider, J., Sachs, T., Giebels, M., Nykänen, H., Shurpali, N.J., Martikainen, P.J., Alm, J., ~~and~~ Wilmking, M.: ~~2007~~. CO₂ flux determination by closed-chamber methods can be seriously biased by inappropriate application of linear regression. *Biogeosciences* 4, 1005–1025. <https://doi.org/10.5194/bg-4-1005-2007>, ~~2007~~.
- Laine, A., Wilson, D., Kiely, G., ~~and~~ Byrne, K.A.: ~~2007~~. Methane flux dynamics in an Irish lowland blanket bog. *Plant Soil* 299, 181–193. <https://doi.org/10.1007/s11104-007-9374-6>, ~~2007~~.

- Livingston, G.P. ~~and~~; Hutchinson, G.L.; ~~1995~~: Enclosure-based measurement of trace gas exchange: applications and sources of error., in: ~~Matson, P.A., Harriss, R.C. (Eds.)~~; Biogenic Trace Gases: Measuring Emissions from Soil and Water, ~~edited by: Matson, P.A., Harriss, R.C.~~; Blackwell Science Ltd., Oxford, UK, pp. 14–51, ~~1995~~.
- Macreadie, P.I., Anton, A., Raven, J.A., Beaumont, N., Connolly, R.M., Friess, D.A., Kelleway, J.J., Kennedy, H., Kuwae, T., Lavery, P.S., Lovelock, C.E., Smale, D.A., Apostolaki, E.T., Atwood, T.B., Baldock, J., Bianchi, T.S., Chmura, G.L., Eyre, B.D., Fourqurean, J.W., Hall-Spencer, J.M., Huxham, M., Hendriks, I.E., Krause-Jensen, D., Laffoley, D., Luisetti, T., Marbà, N., Masque, P., McGlathery, K.J., Megonigal, J.P., Murdiyarso, D., Russell, B.D., Santos, R., Serrano, O., Silliman, B.R., Watanabe, K., ~~and~~ Duarte, C.M.; ~~2019~~: The future of Blue Carbon science.; Nat Commun 10, 3998.; <https://doi.org/10.1038/s41467-019-11693-w>, ~~2019~~.
- Maier, M., Weber, T.K.D., Fiedler, J., Fuß, R., Glatzel, S., Huth, V., Jordan, S., Jurasinski, G., Kutzbach, L., Schäfer, K., Weymann, D., ~~and~~ Hagemann, U.; ~~2022~~: Introduction of a guideline for measurements of greenhouse gas fluxes from soils using non-steady-state chambers.; J. Plant Nutr. Soil Sci. 185, 447–461.; <https://doi.org/10.1002/jpln.202200199>, ~~2022~~.
- Melton, J.R., Wania, R., Hodson, E.L., Poulter, B., Ringeval, B., Spahni, R., Bohn, T., Avis, C.A., Beerling, D.J., Chen, G., Eliseev, A.V., Denisov, S.N., Hopcroft, P.O., Lettenmaier, D.P., Riley, W.J., Singarayer, J.S., Subin, Z.M., Tian, H., Zürcher, S., Brovkin, V., Van Bodegom, P.M., Kleinen, T., Yu, Z.C., ~~and~~ Kaplan, J.O.; ~~2013~~: Present state of global wetland extent and wetland methane modelling: conclusions from a model inter-comparison project (WETCHIMP); Biogeosciences 10, 753–788.; <https://doi.org/10.5194/bg-10-753-2013>, ~~2013~~.
- Moore, T.R. ~~and~~; Knowles, R.; ~~1990~~: Methane emissions from fen, bog and swamp peatlands in Quebec.; Biogeochemistry 11.; <https://doi.org/10.1007/BF00000851>, ~~1990~~.
- Morgan, M.G.; ~~2014~~: Use (and abuse) of expert elicitation in support of decision making for public policy.; Proc. Natl. Acad. Sci. U.S.A. 111, 7176–7184.; <https://doi.org/10.1073/pnas.1319946111>, ~~2014~~.
- Mosier, A.R., Delgado, J.A., Cochran, V.L., Valentine, D.W., ~~and~~ Parton, W.J.; ~~1997~~: Impact of agriculture on soil consumption of atmospheric CH₄ and a comparison of CH₄ and N₂O flux in subarctic, temperate and tropical grasslands.; Nutrient Cycling in Agroecosystems 49, 71–83.; <https://doi.org/10.1023/A:1009754207548>, ~~1997~~.
- Neubauer, S.C.; ~~2021~~: Global Warming Potential Is Not an Ecosystem Property.; Ecosystems 24, 2079–2089.; <https://doi.org/10.1007/s10021-021-00631-x>, ~~2021~~.
- Nickerson, N.R.; ~~2016~~: Evaluating gas emission measurements using Minimum Detectable Flux (MDF); <https://doi.org/10.13140/RG.2.1.4149.2089>, ~~2016~~.
- Pedersen, A.R., Petersen, S.O., ~~and~~ Schelde, K.; ~~2010~~: A comprehensive approach to soil-atmosphere trace-gas flux estimation with static chambers.; European Journal of Soil Science 61, 888–902.; <https://doi.org/10.1111/j.1365-2389.2010.01291.x>, ~~2010~~.
- Pihlatie, M.K., Christiansen, J.R., Aaltonen, H., Korhonen, J.F.J., Nordbo, A., Rasilo, T., Benanti, G., Giebels, M., Helmy, M., Sheehy, J., Jones, S., Juszczak, R., Klefoth, R., Lobo-do-Vale, R., Rosa, A.P., Schreiber, P., Serça, D., Vicca, S., Wolf, B., ~~and~~ Pumpanen, J.; ~~2013~~: Comparison of static chambers to measure CH₄ emissions from soils.; Agricultural and Forest Meteorology 171–172, 124–136.; <https://doi.org/10.1016/j.agrformet.2012.11.008>, ~~2013~~.
- Pirk, N., Mastepanov, M., Parmentier, F.-J.W., Lund, M., Crill, P., ~~and~~ Christensen, T.R.; ~~2016~~: Calculations of automatic chamber flux measurements of methane and carbon dioxide using short time series of concentrations.; Biogeosciences 13, 903–912.; <https://doi.org/10.5194/bg-13-903-2016>, ~~2016~~.
- Pumpanen, J., Kolari, P., Ilvesniemi, H., Minkinen, K., Vesala, T., Niinistö, S., Lohila, A., Larmola, T., Morero, M., Pihlatie, M., Janssens, I., Yuste, J.C., Grünzweig, J.M., Reth, S., Subke, J.-A., Savage, K., Kutsch, W., Østreng, G., Ziegler, W., Anthoni, P., Lindroth, A., ~~and~~ Hari, P.; ~~2004~~: Comparison of different chamber techniques for measuring soil CO₂ efflux.; Agricultural and Forest Meteorology 123, 159–176.; <https://doi.org/10.1016/j.agrformet.2003.12.001>, ~~2004~~.
- Rheault, K., Christiansen, J.R., ~~and~~ Larsen, K.S.; ~~2024~~: goFlux: A user-friendly way to calculate GHG fluxes yourself, regardless of user experience.; JOSS 9, 6393.; <https://doi.org/10.21105/joss.06393>, ~~2024~~.
- Rochette, P. ~~and~~; Eriksen-Hamel, N.S.; ~~2008~~: Chamber Measurements of Soil Nitrous Oxide Flux: Are Absolute Values Reliable? Soil Science Soc of Amer J 72, 331–342.; <https://doi.org/10.2136/sssaj2007.0215>, ~~2008~~.

Rosentreter, J.A., Alcott, L., Maavara, T., Sun, X., Zhou, Y., Planavsky, N.J., ~~and~~ Raymond, P.A., ~~2024~~: Revisiting the Global Methane Cycle Through Expert Opinion. *Earth's Future* 12, e2023EF004234. <https://doi.org/10.1029/2023EF004234>, ~~2024~~.

Saunois, M., Stavert, A.R., Poulter, B., Bousquet, P., Canadell, J.G., Jackson, R.B., Raymond, P.A., Dlugokencky, E.J., Houweling, S., Patra, P.K., Ciais, P., Arora, V.K., Bastviken, D., Bergamaschi, P., Blake, D.R., Brailsford, G., Bruhwiler, L., Carlson, K.M., Carrol, M., Castaldi, S., Chandra, N., Crevoisier, C., Crill, P.M., Covey, K., Curry, C.L., Etiope, G., Frankenberg, C., Gedney, N., Hegglin, M.I., Höglund-Isaksson, L., Hugelius, G., Ishizawa, M., Ito, A., Janssens-Maenhout, G., Jensen, K.M., Joos, F., Kleinen, T., Krummel, P.B., Langenfelds, R.L., Laruelle, G.G., Liu, L., Machida, T., Maksyutov, S., McDonald, K.C., McNorton, J., Miller, P.A., Melton, J.R., Morino, I., Müller, J., Murguía-Flores, F., Naik, V., Niwa, Y., Noce, S., O'Doherty, S., Parker, R.J., Peng, C., Peng, S., Peters, G.P., Prigent, C., Prinn, R., Ramonet, M., Regnier, P., Riley, W.J., Rosentreter, J.A., Segers, A., Simpson, I.J., Shi, H., Smith, S.J., Steele, L.P., Thornton, B.F., Tian, H., Tohjima, Y., Tubiello, F.N., Tsuruta, A., Viovy, N., Voulgarakis, A., Weber, T.S., Van Weele, M., Van Der Werf, G.R., Weiss, R.F., Worthy, D., Wunch, D., Yin, Y., Yoshida, Y., Zhang, W., Zhang, Z., Zhao, Y., Zheng, B., Zhu, Qing, Zhu, Qiu, ~~and~~ Zhuang, Q., ~~2020~~: The Global Methane Budget 2000–2017. *Earth Syst. Sci. Data* 12, 1561–1623. <https://doi.org/10.5194/essd-12-1561-2020>, ~~2020~~.

Schuur, E.A.G., Abbott, B.W., Bowden, W.B., Brovkin, V., Camill, P., Canadell, J.G., Chanton, J.P., Chapin, F.S., Christensen, T.R., Ciais, P., Crosby, B.T., Czimczik, C.I., Grosse, G., Harden, J., Hayes, D.J., Hugelius, G., Jastrow, J.D., Jones, J.B., Kleinen, T., Koven, C.D., Krinner, G., Kuhry, P., Lawrence, D.M., McGuire, A.D., Natali, S.M., O'Donnell, J.A., Ping, C.L., Riley, W.J., Rinke, A., Romanovsky, V.E., Sannel, A.B.K., Schädel, C., Schaefer, K., Sky, J., Subin, Z.M., Tarnocai, C., Turetsky, M.R., Waldrop, M.P., Walter Anthony, K.M., Wickland, K.P., Wilson, C.J., ~~and~~ Zimov, S.A., ~~2013~~: Expert assessment of vulnerability of permafrost carbon to climate change. *Climatic Change* 119, 359–374. <https://doi.org/10.1007/s10584-013-0730-7>, ~~2013~~.

Strack, M., Kellner, E., Waddington, J.M., ~~2005~~: Dynamics of biogenic gas bubbles in peat and their effects on peatland biogeochemistry. *Global Biogeochemical Cycles* 19, 2004GB002330. <https://doi.org/10.1029/2004GB002330>, ~~2005~~.

Treat, C.C., Bloom, A.A., ~~and~~ Marushchak, M.E., ~~2018~~: Nongrowing season methane emissions—a significant component of annual emissions across northern ecosystems. *Global Change Biology* 24, 3331–3343. <https://doi.org/10.1111/gcb.14137>, ~~2018~~.

Virkkala, A.-M., Niittynen, P., Kempainen, J., Marushchak, M.E., Voigt, C., Hensgens, G., Kerttula, J., Happonen, K., Tyystjärvi, V., Biasi, C., Hultman, J., Rinne, J., ~~and~~ Luoto, M., ~~2024~~: High-resolution spatial patterns and drivers of terrestrial ecosystem carbon dioxide, methane, and nitrous oxide fluxes in the tundra. *Biogeosciences* 21, 335–355. <https://doi.org/10.5194/bg-21-335-2024>, ~~2024~~.

Voigt, C., Virkkala, A.-M., Hould Gosselin, G., Bennett, K.A., Black, T.A., Detto, M., Chevrier-Dion, C., Guggenberger, G., Hashmi, W., Kohl, L., Kou, D., Marquis, C., Marsh, P., Marushchak, M.E., Nesic, Z., Nykänen, H., Saarela, T., Sauheitl, L., Walker, B., Weiss, N., Wilcox, E.J., ~~and~~ Sonnentag, O., ~~2023~~: Arctic soil methane sink increases with drier conditions and higher ecosystem respiration. *Nat. Clim. Chang.* 13, 1095–1104. <https://doi.org/10.1038/s41558-023-01785-3>, ~~2023~~.

Waddington, J.M. ~~and~~ Roulet, N.T., ~~1996~~: Atmosphere-wetland carbon exchanges: Scale dependency of CO₂ and CH₄ exchange on the developmental topography of a peatland. *Global Biogeochemical Cycles* 10, 233–245. <https://doi.org/10.1029/95GB03871>, ~~1996~~.

Xu, L., Furtaw, M.D., Madsen, R.A., Garcia, R.L., Anderson, D.J., ~~and~~ McDermitt, D.K., ~~2006~~: On maintaining pressure equilibrium between a soil CO₂ flux chamber and the ambient air. *J. Geophys. Res.* 111, 2005JD006435. <https://doi.org/10.1029/2005JD006435>, ~~2006~~.

Formatted: English (United States)