

RESEARCH ARTICLE

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Key Points:

- We present a $\delta^{13}\text{C}\text{-CH}_4$ record of the rapid CH_4 variation 85ka before present
- We apply Keeling plot analysis and validate the results against a box model
- We propose that tropical wetland emissions caused the observed CH_4 anomaly

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Carbon isotope ratios suggest no additional methane from boreal wetlands during the rapid Greenland Interstadial 21.2

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Abstract Samples from two Greenland ice cores (NEEM and NGRIP) have been measured for methane carbon isotope ratios ($\delta^{13}\text{C}\text{-CH}_4$) to investigate the CH_4 mixing ratio anomaly during Greenland Interstadial (GI) 21.2 (85,000 years before present). This extraordinarily rapid event occurred within 150 years, comprising a CH_4 mixing ratio pulse of 150 ppb ($\sim 25\%$). Our new measurements disclose a concomitant shift in $\delta^{13}\text{C}\text{-CH}_4$ of 1‰. Keeling plot analyses reveal the $\delta^{13}\text{C}$ of the additional CH_4 source constituting the CH_4 anomaly as $-56.8 \pm 2.8\%$, which we confirm by means of a previously published box model. We propose tropical wetlands as the most probable additional CH_4 source during GI-21.2 and present independent evidence that suggests that tropical wetlands in South America and Asia have played a key role. We find no evidence that boreal CH_4 sources, such as permafrost degradation, contributed significantly to the atmospheric CH_4 increase, despite the pronounced warming in the Northern Hemisphere during GI-21.2.

1. Introduction

Methane (CH_4) is a strong greenhouse gas that is accumulating in Earth's atmosphere due to human activity. Currently, CH_4 contributes 20% of the anthropogenic increase in radiative forcing since 1750 [e.g., Forster *et al.*, 2007] and plays a significant role in recent and projected variations of global temperature, sea level, and sea ice extent [e.g., Meehl *et al.*, 2007]. Reconstructions of atmospheric CH_4 mixing ratios (mixing ratios of all gases are indicated by [...] in the following) from ice core samples show distinct variations from decadal [e.g., Grachev *et al.*, 2007; Chappellaz *et al.*, 2013; Mitchell *et al.*, 2013] to orbital timescales [e.g., Loulergue *et al.*, 2008]. Rapid [CH_4] increases of the order of 150–300 ppb occurred over decades to centuries in association with Northern Hemispheric warm events [e.g., Brook *et al.*, 1996; Grachev *et al.*, 2007]. We use the most recent nomenclature after Rasmussen *et al.* [2014] and name the cold periods "Greenland Stadial" (GS) and the intermittent warm events "Greenland Interstadial" (GI).

High-resolution data sets show that some GI events were preceded by sharp precursor events of high amplitude, e.g., GI-21.2 [e.g., Grachev *et al.*, 2007; Capron *et al.*, 2010; Boch *et al.*, 2011; Vallelonga *et al.*, 2012; Chappellaz *et al.*, 2013; Deplazes *et al.*, 2013]. The rapid GI-21.2 event about 85,000 years before 2000 A.D. (b2k) comprised a [CH_4] spike of ~ 150 ppb that occurred within ~ 150 years (Figure 1). This extraordinary event is marked by the highest [CH_4] growth rate recorded in Greenland ice cores [Chappellaz *et al.*, 2013]. Rapid [CH_4] changes of this magnitude and timescale are of particular interest for studies on the biogeochemistry of CH_4 and the sensitivity of CH_4 source fluxes to climate change.

Atmospheric chemistry models suggest that the [CH_4] variability during GI events was mostly driven by variations in the sources of CH_4 rather than the sinks [Levine *et al.*, 2012]. Atmospheric CH_4 sources have distinct isotopic compositions in both $\delta^{13}\text{C}$ and $\delta^2\text{H}$ that depend on the source processes and the CH_4 precursor material [e.g., Quay *et al.*, 1999; Whiticar and Schaefer, 2007]. Observations of $\delta^{13}\text{C}\text{-CH}_4$ or $\delta^2\text{H}\text{-CH}_4$ can therefore constrain CH_4 source flux reconstructions, which can then be interpreted in the context of climate variability [e.g., Ferretti *et al.*, 2005; Schaefer *et al.*, 2006; Sowers, 2006; Fischer *et al.*, 2008; Bock *et al.*, 2010; Sapart *et al.*, 2012].

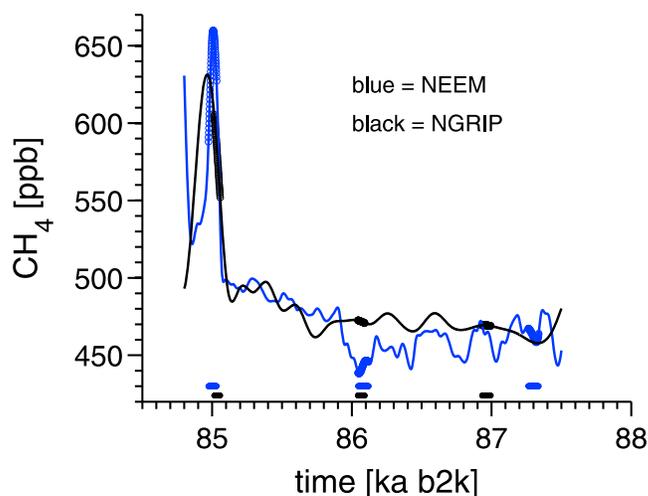


Figure 1. $[\text{CH}_4]$ spline fits for NEEM (blue) and NGRIP (black). Circles on the lines and bars at the bottom of the plot highlight the locations of the $\delta^{13}\text{C}-\text{CH}_4$ samples relative to $[\text{CH}_4]$.

Isotope records of CH_4 suggest tropical and/or boreal wetland source flux variations as main drivers of the $[\text{CH}_4]$ variability [e.g., Sowers, 2006; Fischer *et al.*, 2008; Mischler *et al.*, 2009; Bock *et al.*, 2010; Möller *et al.*, 2013]. Furthermore, isotopic evidence clearly demonstrates that CH_4 hydrate destabilization was not the primary CH_4 source that caused the $[\text{CH}_4]$ variability during the last deglaciation [Sowers, 2006], GI-7 and GI-8 [Bock *et al.*, 2010].

Most studies that analyzed the isotopic variability of CH_4 in ice core samples use mass balance calculations in box models [e.g., Ferretti *et al.*, 2005; Schaefer *et al.*, 2006; Sowers, 2006; Fischer *et al.*, 2008; Mischler *et al.*, 2009; Bock *et al.*, 2010; Sapart *et al.*, 2012]. Here we investigate the change in $\delta^{13}\text{C}-\text{CH}_4$ that is associated with the $[\text{CH}_4]$ anomaly of GI-21.2 using Keeling plot analysis (KPA) [Keeling, 1958]. KPA is technically a two-component mixing model that provides the $\delta^{13}\text{C}-\text{CH}_4$ of an additional CH_4 source, which in our case is the source of the $[\text{CH}_4]$ anomaly of GI-21.2. We present a Monte Carlo technique that considers both the analytical and data processing errors as well as a potential sampling bias in order to estimate the uncertainty of the KPA. Then, we compare the $\delta^{13}\text{C}-\text{CH}_4$ result from the KPA to the result from the forward stepping box model of Lassey *et al.* [2007] and show that the two methods agree well within the uncertainty of the KPA.

Recently, Möller *et al.* [2013] suggested that $\delta^{13}\text{C}-\text{CH}_4$ and $[\text{CH}_4]$ vary independently on millennial to glacial-interglacial timescales, which also questions the suitability of mass balance calculations for CH_4 source reconstructions on the timescales of our study period. We therefore test alternative scenarios where the $\delta^{13}\text{C}-\text{CH}_4$ excursion of GI-21.2 is superimposed on a long-term $\delta^{13}\text{C}-\text{CH}_4$ trend that is controlled by $[\text{CO}_2]$. We argue that KPA and mass balance calculations can be used in our study period when the effect of possible $\delta^{13}\text{C}-\text{CH}_4$ background scenarios is carefully considered.

To our knowledge, this is the first time KPA is applied to studies of CH_4 in ice core samples. Therefore, we first review assumptions and necessary conditions for the use of KPA and then explain the processes that need to be considered in order to reconstruct the $\delta^{13}\text{C}$ of an additional CH_4 source from ice core samples. Finally, we discuss and evaluate CH_4 emission scenarios in the context of other, independent climate records.

2. Methods

2.1. Measurement Techniques

Our analytical technique for $\delta^{13}\text{C}-\text{CH}_4$ analysis is described in detail by Sperlich *et al.* [2013]. In short, cleaned ice core samples are melted in a vacuum system from which the liberated air sample is extracted. A helium carrier gas stream transports the sample through the analytical system to isolate CH_4 and to combust it into CO_2 before it is measured for $\delta^{13}\text{C}$ on an isotope ratio mass spectrometer. Our $\delta^{13}\text{C}$ results are reported on the Vienna Pee Dee Belemnite isotope scale, using the referencing technique described by Sperlich *et al.* [2012]. The analytical uncertainty of the $\delta^{13}\text{C}-\text{CH}_4$ measurements is 0.09‰. Our analytical method is free of krypton artifacts which have recently been identified as a major problem in $\delta^{13}\text{C}-\text{CH}_4$ measurements [Schmitt *et al.*, 2013].

Optical measurement techniques were recently applied to measure $[\text{CH}_4]$ in ice core samples [Stowasser *et al.*, 2012] and provided a continuous record of unprecedented temporal resolution and precision from the NEEM ice core [Chappellaz *et al.*, 2013]. This new method presents the $[\text{CH}_4]$ of GI-21.2 in unprecedented detail. Discrete $[\text{CH}_4]$ measurements from the NGRIP ice core show the anomaly of GI-21.2 in comparable magnitude [Baumgartner *et al.*, 2014]. We fitted two splines to the $[\text{CH}_4]$ records from Chappellaz *et al.* [2013] and Baumgartner *et al.* [2014] to calculate the mean $[\text{CH}_4]$ of our $\delta^{13}\text{C}\text{-CH}_4$ samples (Figure 1). The spline of the NEEM data includes a 25 year smoothing filter, while replicate measurements of $[\text{CH}_4]$ in the NGRIP record were averaged to avoid artifacts. The NEEM and the NGRIP spline fit show a slightly different timing for GI-21.2. However, timing offsets between the records can be corrected for because $[\text{CH}_4]$ varies synchronously in the atmosphere [Blunier *et al.*, 2007]. The higher temporal resolution of the NEEM $[\text{CH}_4]$ record justifies to transfer the NGRIP $[\text{CH}_4]$ data to the NEEM timescale for GI-21.2. Therefore, we used the NEEM spline in our box model calculations (section 2.6).

2.2. Ice Core Samples for $\delta^{13}\text{C}\text{-CH}_4$ Measurements

The rapid GI-21.2 event is recorded within only ~ 1.5 m in the NEEM ice core [Chappellaz *et al.*, 2013]. Because the number of available ice core samples from GI-21.2 and GS-22 is extremely limited (0.55 m ice per sample), we used samples from both the NEEM and NGRIP ice cores for $\delta^{13}\text{C}\text{-CH}_4$ analysis. We measured four samples of the stadial period preceding GI-21.2 (GI-22), when $[\text{CH}_4]$ was stable. We analyzed two samples of GI-21.2 and six younger samples of the strong $[\text{CH}_4]$ variation of GI-21.1e (Figure 2). The age interval that is integrated within each of our $\delta^{13}\text{C}\text{-CH}_4$ samples from GS-22 and GI-21.2 is displayed in relation to the $[\text{CH}_4]$ history in Figure 1. Together, our two GI-21.2 samples integrate more than 50% of the event, including the $[\text{CH}_4]$ peak and the highest rates of $[\text{CH}_4]$ change.

2.3. Applied Corrections on $\delta^{13}\text{C}\text{-CH}_4$ Measurements

All $\delta^{13}\text{C}\text{-CH}_4$ measurements are corrected for firn diffusion fractionation after Buizert *et al.* [2013]. This correction depends on the physical properties of the respective gas species, its relative growth rate, and a site-specific time-dependent factor, which is determined by the diffusive column height. The firn diffusion fractionation is generally smaller at Greenlandic than at central Antarctic sites. Our firn diffusion correction reaches maximum values of 0.34‰ during GI-21.2. This semiempirical method is published with a general, relative uncertainty of 30% for Greenland ice cores.

We also correct the $\delta^{13}\text{C}\text{-CH}_4$ and $[\text{CH}_4]$ data for the disequilibrium effect [Tans, 1997] that we determined with the box model of Lassey *et al.* [2007]. The disequilibrium effect for both $\delta^{13}\text{C}\text{-CH}_4$ and $[\text{CH}_4]$ is most pronounced for our GI-21.2 samples, where it accounts for up to 0.12‰ and 24 ppb, respectively. We discuss the relevance of this correction on our KPA results in section 3.1.

2.4. Assumptions for the Analysis of $\delta^{13}\text{C}\text{-CH}_4$ During Greenland Interstadial 21.2 Using Mass Balance Calculations

Because the limited amount of sample ice restricts the temporal data resolution, even our high-resolution record with an average resolution of one sample per 380 years includes a data gap of ~ 1000 years, just before the onset of the GI-21.2 event. This requires an assumption as to the timing of the $\delta^{13}\text{C}\text{-CH}_4$ variation before the GI-21.2 event, which is critical for the analysis.

We can think of three different $\delta^{13}\text{C}\text{-CH}_4$ background scenarios:

1. The variation of $\delta^{13}\text{C}\text{-CH}_4$ is correlated with $[\text{CH}_4]$; i.e., the $\delta^{13}\text{C}\text{-CH}_4$ background does not change before the onset of GI-21.2 (Figure 2c). This scenario is in agreement with most of the existing publications on $\delta^{13}\text{C}\text{-CH}_4$ in ice core samples [e.g., Ferretti *et al.*, 2005; Schaefer *et al.*, 2006; Sowers, 2006; Fischer *et al.*, 2008; Mischler *et al.*, 2009; Bock *et al.*, 2010; Sapart *et al.*, 2012]. In particular, Melton *et al.* [2012] recently reported a variation in $\delta^{13}\text{C}\text{-CH}_4$ in correspondence to a rapid change in $[\text{CH}_4]$. In the following, we will refer to this as the $[\text{CH}_4]$ -correlated background scenario.
2. The $\delta^{13}\text{C}\text{-CH}_4$ is controlled by $[\text{CO}_2]$ on millennial to glacial timescales (Figure 2b) as recently published by Möller *et al.* [2013]. We will name this the $[\text{CO}_2]$ -correlated background scenario throughout this study.
3. A $\delta^{13}\text{C}\text{-CH}_4$ depletion that is continuous from the last measured pre-event value to the peak of GI-21.2 (Figure 2a).

Other evolutions of the $\delta^{13}\text{C}\text{-CH}_4$ background cannot be ruled out but are highly speculative. We reject scenario 3 (as well as other $\delta^{13}\text{C}\text{-CH}_4$ background histories), because there is no mechanistic explanation for a $\delta^{13}\text{C}\text{-CH}_4$ variability that is not linked to either $[\text{CH}_4]$ or $[\text{CO}_2]$.

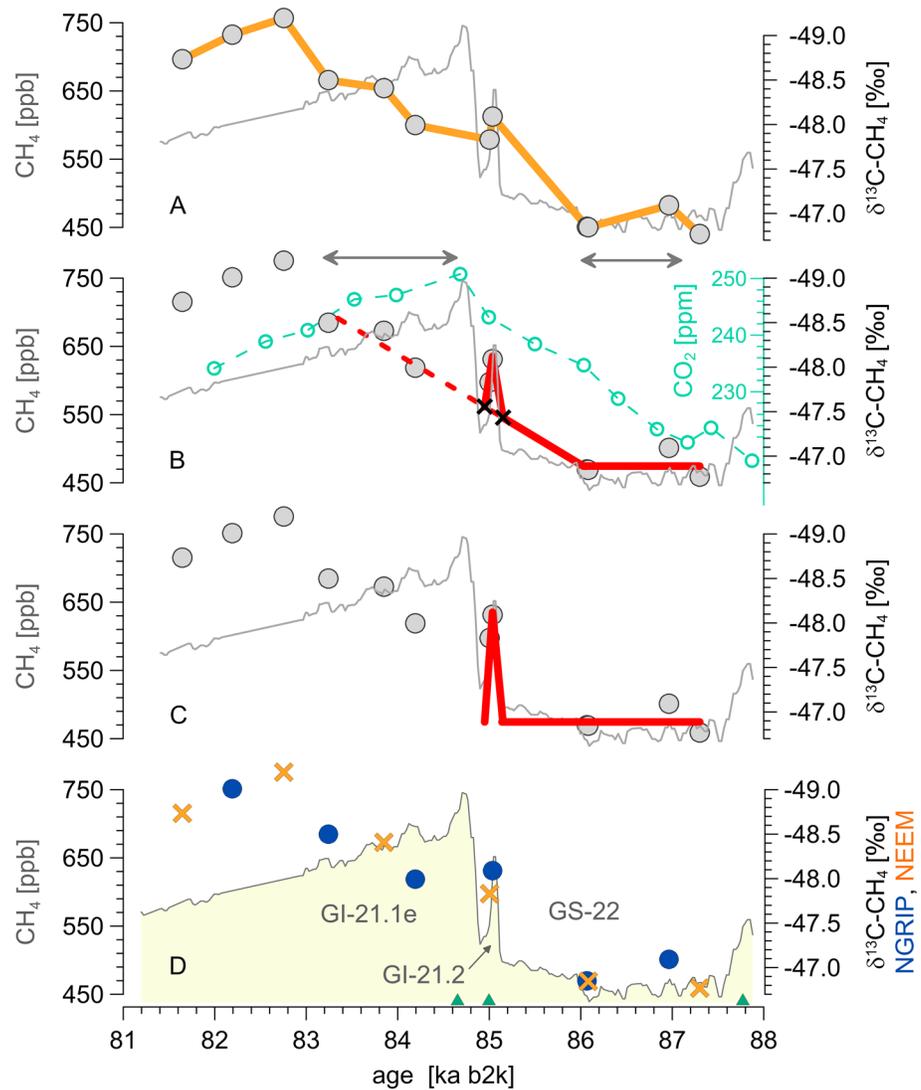


Figure 2. $\delta^{13}\text{C}\text{-CH}_4$ background scenarios: (left y axes) $[\text{CH}_4]$, grey lines [Chappellaz et al., 2013]; (right y axes) $\delta^{13}\text{C}\text{-CH}_4$, filled symbols. Figures 2a–2c show the discussed scenarios of the $\delta^{13}\text{C}\text{-CH}_4$ background during the sample gap between 85–86 ka b2k and the GI-21.2 event. (a) The $\delta^{13}\text{C}\text{-CH}_4$ as direct (orange) line between measurements. (b) $[\text{CO}_2]$ [Bereiter et al., 2012], $\delta^{13}\text{C}\text{-CH}_4$ reconstruction based on the $[\text{CO}_2]$ -correlated background (red line) as determined by the regression (red, dashed line); two arrows indicate the time lag between $[\text{CO}_2]$ and $\delta^{13}\text{C}\text{-CH}_4$ that averages to ~1270 years. (c) $\delta^{13}\text{C}\text{-CH}_4$ reconstruction based on the $[\text{CH}_4]$ -correlated background scenario (red line). (d) The $\delta^{13}\text{C}\text{-CH}_4$ measurements from NGRIP and NEEM ice cores. Note the good agreement between the 86 ka samples from both ice cores and the ^{13}C enrichment with decreasing $[\text{CH}_4]$ at the end of the GI-21.2 event. Green triangles (Figure 2d) indicate points to match $[\text{CH}_4]$ from the EPICA Dronning Maud Land (EDML) ice core [Schilt et al., 2010] and NEEM to transfer $[\text{CO}_2]$ from the EDML gas age scale to GICC05_modelext [Blunier et al., 2007].

Also, the $\delta^{13}\text{C}\text{-CH}_4$ peak and decline of GI-21.2 indicate that the $\delta^{13}\text{C}\text{-CH}_4$ signal is indeed a positive excursion and not just part of a longer-term trend. Changes in $[\text{CH}_4]$ indicate variations in CH_4 emissions (assuming no or small sink variability). These necessarily incur changes in $\delta^{13}\text{C}\text{-CH}_4$ of the total source and consequently the atmosphere, except for the unlikely case that all sources change by the same relative amounts. Therefore, the coinciding excursions in $[\text{CH}_4]$ and $\delta^{13}\text{C}\text{-CH}_4$ can be analyzed for the underlying source changes [Ferretti et al., 2005; Schaefer et al., 2006; Sowers, 2006; Fischer et al., 2008; Mischler et al., 2009; Bock et al., 2010; Sapart et al., 2012]. The $[\text{CO}_2]$ -correlated $\delta^{13}\text{C}\text{-CH}_4$ variability [Möller et al., 2013] must also be taken into account. However, this can only explain a part of the GI-21.2 $\delta^{13}\text{C}\text{-CH}_4$ excursion. A significant $\delta^{13}\text{C}\text{-CH}_4$ deviation linked to the $[\text{CH}_4]$ peak is evident even if superimposed on a longer-term, $[\text{CO}_2]$ -correlated trend in $\delta^{13}\text{C}\text{-CH}_4$ (Figure 2b). Therefore, we hypothesize that scenarios 1 and 2 represent the outer bounds for the possible

evolution of the $\delta^{13}\text{C}-\text{CH}_4$ background, in agreement with previous publications. The $\delta^{13}\text{C}-\text{CH}_4$ excursion that is superimposed on the background trends can be analyzed with KPA and mass balance calculations because of its short duration of GI-21.2 and the likelihood of little variability in $\delta^{13}\text{C}-\text{CH}_4$ background. We present separate solutions for scenarios 1 and 2. Because there is no evidence to prefer either the $[\text{CH}_4]$ - or the $[\text{CO}_2]$ -correlated background scenario, we use the average of both analyses and their propagate uncertainty.

Furthermore, we argue that our measurements and samples are appropriate to reconstruct the atmospheric variability of GI-21.2. Air is subject to mixing and diffusion processes in the firn column, before air is permanently trapped in the ice [e.g., Buizert *et al.*, 2012], which smoothes the recorded atmospheric signal. Ice core studies are therefore liable to underestimate atmospheric variability. However, previous studies showed excellent agreement between overlapping $\delta^{13}\text{C}-\text{CH}_4$ measurements in atmospheric, firn-air, and ice core samples [Francey *et al.*, 1999; Ferretti *et al.*, 2005] (note that potential disagreements between multiple $\delta^{13}\text{C}-\text{CH}_4$ firn-air records are largely based on analytical artifacts and laboratory offsets [Sapart *et al.*, 2013] that may range in the order of 0.5‰ [Sapart *et al.*, 2011]). The ice core samples from Ferretti *et al.* [2005] originated from a high-accumulation site where firn smoothing window is smaller than in central Greenland (≤ 20 years [Ferretti *et al.*, 2005] versus ≤ 80 years with mean of ~ 25 years [Spahni *et al.*, 2003]). Furthermore, the $[\text{CH}_4]$ growth rate was significantly larger during the time period studied by Ferretti *et al.* [2005] compared to that of GI-21.2 (5–17 ppb/yr [Etheridge *et al.*, 1998] versus 2.5 ppb/yr [Chappellaz *et al.*, 2013]), where a larger growth rate increases the impact of firn column effects. Therefore, we expect that firn smoothing has a negligible effect on our record. Our two samples of the GI-21.2 event integrate about 50% of the $[\text{CH}_4]$ variability (Figure 1), which might dampen the atmospheric signal. Because we average $[\text{CH}_4]$ over the exact time period that is integrated in each $\delta^{13}\text{C}-\text{CH}_4$ sample for the KPA, we expect that the impact of the sample integration on the KPA result is not significant.

It is furthermore important to note that our $\delta^{13}\text{C}-\text{CH}_4$ samples integrate 40–70 years between sample top and bottom. For the KPA, we use $[\text{CH}_4]$ and $\delta^{13}\text{C}-\text{CH}_4$ of identical ice cores and average $[\text{CH}_4]$ over the time period that is integrated in each $\delta^{13}\text{C}-\text{CH}_4$ sample. Because of this time integration, our analysis resolves variations as 40–70 year averages.

2.5. Keeling Plot Analysis

2.5.1. Keeling Plot Analysis for CH_4 Source Determination in Atmospheric Samples

For the KPA, the isotopic composition of a group of samples is plotted versus its inverse mixing ratio. The intercept of the linear regression with the y axis indicates the average isotopic signature of a trace gas source [Keeling, 1958]. This technique represents a two-component mixing model with background air and one additional source of analyte gas as the only principal components [Pataki *et al.*, 2003]. Pataki *et al.* [2003] express the global atmospheric budget as

$$c_a = c_b + c_s \quad (1)$$

where c_a , c_b , and c_s represent the CH_4 mixing ratios as measured in the atmosphere, the background atmosphere, and the term due to the additional source, respectively. While c_a and c_b are directly measured, c_s can be calculated after equation (1), because we assume that the background is well defined by the measurements of c_b .

Referring to the isotopic composition of each term by $\delta^{13}\text{C}$, the mass balance calculation

$$\delta^{13}\text{C}_a c_a = \delta^{13}\text{C}_b c_b + \delta^{13}\text{C}_s c_s \quad (2)$$

allows to calculate the $\delta^{13}\text{C}_s$ of the additional CH_4 source that accounts for c_s in the present-day atmosphere [e.g., Fisher *et al.*, 2011].

2.5.2. Keeling Plot Analysis for CH_4 Source Analysis in Ice Core Samples

Unlike for direct atmospheric measurements (e.g., in the vicinity of CH_4 point sources), KPA on $\delta^{13}\text{C}-\text{CH}_4$ in ice core samples does not provide the isotopic composition of an additional CH_4 source directly, because two additional mechanisms have to be taken into account. There are the atmospheric disequilibrium effect [e.g., Tans, 1997] and the impact of the atmospheric sink fractionation (ϵ) on the $\delta^{13}\text{C}$ of the additional CH_4 source. Every ice core measurement needs to be corrected for the average disequilibrium effect per sample, which can be determined with a box model based on measured $[\text{CH}_4]$ and $\delta^{13}\text{C}-\text{CH}_4$ time series, resulting in $\delta^{13}\text{C}_{a-\text{corr}}$, $\delta^{13}\text{C}_{b-\text{corr}}$, $c_{a-\text{corr}}$, and $c_{b-\text{corr}}$. Secondly, the $\delta^{13}\text{C}_s$ term in equation (2) has been altered by atmospheric sink

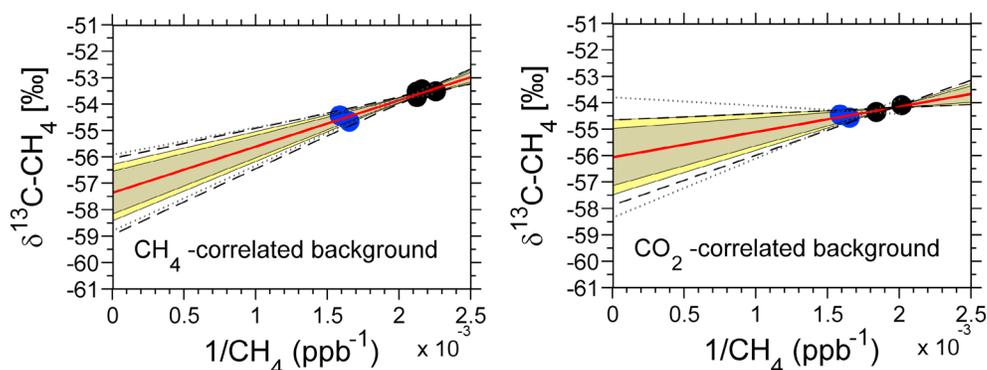


Figure 3. Keeling plots with uncertainties. (left) Keeling Plot with two measurements from GI-21.2 (blue circles) and [CH₄]-correlated background as represented by four measurements from GS-22 (black circles). (right) Keeling Plot of [CO₂]-correlated background, similar to Figure 3 (left) except that the two black circles indicate the two artificial data points. Red lines represent the linear fit; grey and yellow shading illustrate the 95 and 99% confidence intervals of the linear fit as determined by the Monte Carlo analysis, respectively. Dotted lines indicate the least squares uncertainty; dashed lines indicate the uncertainty estimate of the “quasi” bootstrap.

fractionation of a similar magnitude as $\delta^{13}C_a$ and $\delta^{13}C_b$. By solving equation (2) for $\delta^{13}C_s$ and accounting for the disequilibrium effect correction and the sink weighted isotope fractionation (ϵ_{tot}), we get

$$\delta^{13}C_s = \frac{(\delta^{13}C_{a-corr} \times c_{a-corr} - \delta^{13}C_{b-corr} \times c_{b-corr})}{c_s} \times \left(1 + \frac{\epsilon_{tot}}{1000}\right) + \epsilon_{tot} \quad (3)$$

and are able to calculate the $\delta^{13}C$ of the additional CH₄ source in ice core samples.

2.5.3. The Isotopic Fractionation of the Total CH₄ Sink in Keeling Plot Analysis

Literature values of ϵ_{tot} vary from -7.7‰ [Lassey *et al.*, 2007] to -5.4‰ [Mischler *et al.*, 2009], while it has been suggested to scale ϵ_{tot} to [CH₄] on glacial-interglacial timescales [Schaefer and Whiticar, 2008]. The determination of ϵ_{tot} depends on the isotope fractionation of the respective CH₄ sinks and their relative contribution to the total sink fluxes. Note that both factors are likely to have varied on glacial-interglacial timescales and that the cumulative effect on ϵ_{tot} on glacial timescales is not well understood [e.g., Schaefer and Whiticar, 2008; Levine *et al.*, 2012]. The weak correlation between [CH₄] and $\delta^{13}C\text{-CH}_4$ on millennial to glacial timescales [Möller *et al.*, 2013] further suggests that the forcing of [CH₄] on ϵ_{tot} is not controlling $\delta^{13}C\text{-CH}_4$. We apply the ϵ_{tot} of -7.0‰ that Schaefer and Whiticar [2008] calculated for the preindustrial Holocene and discuss the effect of different ϵ_{tot} scenarios on our results in section 3.1.

2.5.4. Uncertainty Estimate of Least Squares Fit in Keeling Plot Analysis

A linear regression is used to calculate the slope and intercept of the KPA. This is often undertaken using a least squares fit, where the confidence interval of the fit is calculated based on the uncertainty of both slope and intercept, the degrees of freedom ($n - 2$), and the according quantiles of a Student's t distribution at the chosen confidence level (Figure 3). We see two disadvantages of the least squares uncertainty in our case:

1. It requires homoscedasticity, whereas the residuals of our linear fit show a bimodal distribution of the variance with clusters during the background period (GS-22) and during GI-21.2. The residuals of GI-21.2 data are larger by a factor of 2. Violating the precondition of homogenous variance of the residuals might therefore produce a data-specific bias, which we cannot investigate further because of the limited number of samples.
2. The least squares uncertainty neither considers the analytical uncertainty of 0.09‰ nor the uncertainty of the firn diffusion correction. The latter accounts for 0.11‰ in both GI-21.2 samples and for $\leq 0.01\text{‰}$ in all samples from GS-22 and GI-21.1e. The combined analytical uncertainty is thus clustered with higher uncertainties during GI-21.2, which is not reflected in the least squares scenario. We address these issues with a Monte Carlo estimate of the uncertainty.

2.5.5. Uncertainty Estimate of Keeling Plot Analysis Using Monte Carlo Technique

We designed a Monte Carlo simulation to determine 95 and 99% confidence intervals for least squares linear regressions based on the propagated uncertainties of each data point. In this approach, we randomly perturbed all data points independently, then performed a regression on the perturbed data, and repeated this process 10,000 times.

We calculated the perturbations as the sum of two perturbations, one based on a Gaussian distribution with a standard deviation equal to the measurement uncertainty (0.09‰) and one with a standard deviation equal to the data point specific uncertainty of the firn diffusion correction (0.01–0.11‰). These two perturbations were calculated independently, such that they might have an additive impact for some iterations and a canceling effect for others. The 95 and 99% confidence intervals were then calculated as the 2.5–97.5 and 0.5–99.5 percentile values of the resulting regression curves. Because we have only two data points from GI-21.2, we constructed a quasi bootstrap method to account for errors from sampling bias. Here the same Monte Carlo routine used the background data of the four GI-22 samples, but it was assigned either one or the other data point of GI-21.2 twice, each time in combination with a new randomized uncertainty. This quasi bootstrap method calculates another two sets of 10,000 regressions for each case, that either of our two GI-21.2 samples represents the true value better than both values together. We show the 95 and 99% confidence intervals of the complete data set and furthermore the uppermost and lowermost 99% confidence interval boundaries of the quasi bootstrap in Figure 3. Note that the range between the latter two compares well to the uncertainties of the least squares method (section 2.5.4). We consider the Monte Carlo approach to be the more robust uncertainty estimate for several reasons: (1) It enables to calculate 95 and 99% of the linear fits that can all be seen as best scenarios within the uncertainty of our data. Therefore, it provides valuable, structural information of the uncertainty range. (2) It is free of assumptions regarding the distribution of the residuals. (3) The quasi bootstrap method quantifies the sampling bias.

2.5.6. Determining the [CO₂]-Correlated $\delta^{13}\text{C-CH}_4$ Background Scenario for Keeling Plot Analysis of GI-21.2

Möller *et al.* [2013] show the surprisingly strong correlation between $\delta^{13}\text{C-CH}_4$ and [CO₂] during the last 160 ka and highlight periods when the correlation seems to break down, including the period after the onset of GI-21.1. The latter occurs during the 2300 year data gap in their record between 80.3 and 82.6 ka b2k Before the sample at 82.6 ka b2k, i.e., during our study period, $\delta^{13}\text{C-CH}_4$ and [CO₂] seem to correlate [Möller *et al.*, 2013, Figure 1]. However, the average temporal resolution of 1660 years in this part of the record from Möller *et al.* [2013] complicates the precise timing of the correlation breakdown. Our data of this period are of higher temporal resolution and show a correlation between the $\delta^{13}\text{C-CH}_4$ background and [CO₂] during GS-22 and the early part of GI-21.1e, if we allow for an average time lag between [CO₂] and $\delta^{13}\text{C-CH}_4$ of ~1270 years (Figure 2b). Interestingly, this time lag is in agreement with Burckel *et al.* [2015], who found that changes in the Atlantic Meridional Overturning Circulation (AMOC) preceded precipitation changes in tropical South America during the last glacial period by 500–1700 years. This could provide an additional mechanistic clue to the correlation between [CO₂] and $\delta^{13}\text{C-CH}_4$ as interpreted by Möller *et al.* [2013], as the AMOC is related to changes in atmospheric [CO₂] [Bereiter *et al.*, 2012] and the precipitation in South America to tropical CH₄ emissions, where the latter impact $\delta^{13}\text{C-CH}_4$ (Table 2). Note that independently of a precise determination of the time lag, the correlation between $\delta^{13}\text{C-CH}_4$ and [CO₂] during the period just before and after GI-21.2 is evident. Therefore, the assumption of the [CO₂]-correlated background scenario with monotonous $\delta^{13}\text{C-CH}_4$ depletion around GI-21.2 is justified.

In order to determine the [CO₂]-correlated $\delta^{13}\text{C-CH}_4$ background scenario, we use three of our measurements from GI-21.1e that are ≤ 2000 years younger than GI-21.2. We fitted a regression through these three data points from GI-21.1e and the latest two measurements of GS-22 (red dashed line in Figure 2b). The resulting [CO₂]-correlated background scenario for GI-21.2 and GS-22 is indicated by the thick red line in Figure 2b. Then, we calculated the $\delta^{13}\text{C-CH}_4$ in two points from just before and just after GI-21.2 based on the regression (black crosses in Figure 2b) and determined the corresponding [CH₄] data from Chappellaz *et al.* [2013] (Table 1). The two generated data pairs simulate a hypothetical background scenario for the case that the [CO₂]-correlated trend in $\delta^{13}\text{C-CH}_4$ represents the atmospheric $\delta^{13}\text{C-CH}_4$ variation around GI-21.2 more accurately than the [CH₄]-correlated background (thick red line in Figure 2c). A KPA was then performed using the two artificial data points as background and our measured data from GI-21.2. This experiment highlights two important results: (1) GI-21.2 shows a significant $\delta^{13}\text{C-CH}_4$ event even when superimposed on a long-term $\delta^{13}\text{C-CH}_4$ trend (Table 1). (2) Using both background scenarios for KPA leads to similar CH₄ source signatures that agree within the uncertainty estimate (Figure 3). In this study, the background assumption does not impact on our interpretation of the KPA result.

2.6. Configuration of a Box Model to Test Results From Keeling Plot Analysis

We configured the box model of Lassey *et al.* [2007] so that a base source and an additional source can be implemented in order to compare KPA and box model outputs. Specifically, the KPA assumes that atmosphere,

Table 1. $\delta^{13}\text{C}\text{-CH}_4$ and $[\text{CH}_4]$ Data Used for Keeling Plot Analysis^a

| Ice Core | Age(ka b2k) | $\delta^{13}\text{C}\text{-CH}_4(\text{‰})$ | $[\text{CH}_4](\text{ppb})$ |
|----------|-------------|---|-----------------------------|
| NEEM | 85.003 | -47.83 | 635 |
| NGRIP | 85.036 | -48.12 | 581 |
| NGRIP | 86.068 | -46.85 | 472 |
| NEEM | 86.083 | -46.84 | 444 |
| NGRIP | 86.964 | -47.09 | 469 |
| NEEM | 87.301 | -46.77 | 463 |
| — | 84.950 | -47.56 ^b | 553 ^c |
| — | 85.150 | -47.43 ^b | 496 ^c |

^aColumns 1–4 show sample site, mean gas age on the GICC05 time scale [Wolff *et al.*, 2010], $\delta^{13}\text{C}\text{-CH}_4$ after correction for firn diffusion fractionation, and mean $[\text{CH}_4]$ per $\delta^{13}\text{C}\text{-CH}_4$ sample, respectively.

^bArtificial $\delta^{13}\text{C}\text{-CH}_4$ values for $[\text{CO}_2]$ -correlated background (Figure 2b).

^c $[\text{CH}_4]$ values for $[\text{CO}_2]$ -correlated background from Chappellaz *et al.* [2013].

sources, and sink are in equilibrium, whereas the box model accounts for disequilibrium effects as discussed by Tans [1997]. The CH_4 source flux in the box model is set to reconstruct a smoothed spline fit (25 year smoothing window) of the NEEM $[\text{CH}_4]$ record from Chappellaz *et al.* [2013] (Figure 4a). The background source is configured to simulate the atmospheric $[\text{CH}_4]$ history during GS-22 by varying the source fluxes of background CH_4 while a constant $\delta^{13}\text{C}$ is assigned (Figure 4d). The $\delta^{13}\text{C}$ of background CH_4 was estimated as -53.6‰ by averaging the $\delta^{13}\text{C}\text{-CH}_4$ of our samples from GS-22 and correcting them for ϵ_{tot} of -7.0‰ (equation (3)). The onset of the GI-21.2 event occurs in the year 85,100 b2k in our $[\text{CH}_4]$ spline fit. The box model is configured to use a constant flux of the background source (fixed at the 85,100 b2k value) thereafter and to match the atmospheric $[\text{CH}_4]$ history of GI-21.2 by varying the flux of the additional CH_4 source (Figure 4e) with a $\delta^{13}\text{C}\text{-CH}_4$ that can be varied. The $\delta^{13}\text{C}\text{-CH}_4$ value of the additional source is chosen so that the atmospheric $\delta^{13}\text{C}\text{-CH}_4$ history is in best agreement with our two $\delta^{13}\text{C}\text{-CH}_4$ measurements from GI-21.2 (red circles in Figure 4e). We defined best agreement such that the difference to the modeled $\delta^{13}\text{C}\text{-CH}_4$ history is identical for both of our $\delta^{13}\text{C}\text{-CH}_4$ samples. The box model solves for the $\delta^{13}\text{C}$ of the additional CH_4 during GI-21.2, which we compare to the $\delta^{13}\text{C}\text{-CH}_4$ result of the KPA. We also apply the $\delta^{13}\text{C}\text{-CH}_4$ background of the $[\text{CO}_2]$ -correlated scenario in a box model run to compare the results to the KPA.

In addition, the box model is used to estimate the disequilibrium effect that results from changes in CH_4 source fluxes and the $\delta^{13}\text{C}$ of CH_4 sources [Tans, 1997]. The impact of the disequilibrium effect on $\delta^{13}\text{C}$ scales with ϵ_{tot} , where a stronger sink fractionation causes a stronger disequilibrium. The disequilibrium effect variation during GS-22 and GI-21.2 is shown in Figures 4b and 4c. It is used to correct both the $\delta^{13}\text{C}$ and the mean $[\text{CH}_4]$ of our $\delta^{13}\text{C}\text{-CH}_4$ samples for the KPA (equation (3)) and has a small impact on our KPA result.

3. Results

3.1. Results From Keeling Plot Analysis

Our $\delta^{13}\text{C}\text{-CH}_4$ measurements, the corresponding $[\text{CH}_4]$ data, and the two pairs of artificial data for the $[\text{CO}_2]$ -correlated background scenario are shown in Table 1. The KPA with $[\text{CH}_4]$ -correlated $\delta^{13}\text{C}\text{-CH}_4$ background intersects the y axis at -57.4‰ (Figure 3, left). We derive a conservative uncertainty estimate of $\pm 2\text{‰}$ from the 99% interval of the quasi bootstrap technique (section 2.5.5). The KPA for the $[\text{CO}_2]$ -correlated background scenario intersects the y axis at $-56.1 \pm 2\text{‰}$ (Figure 3, right). The difference between the two KPA results is 1.3‰ , which is well within the uncertainty. The good agreement between KPA using both background scenarios suggests that the assumption regarding the background variation of $\delta^{13}\text{C}\text{-CH}_4$ during GS-22 is not critical for our interpretation of the results.

Applying the ϵ_{tot} of -5.4‰ from Mischler *et al.* [2009] to our analysis would shift the KPA results by 1.5‰ toward more enriched $\delta^{13}\text{C}$ values. Scaling ϵ_{tot} to $[\text{CH}_4]$ due to changes in the soil sink flux as suggested by Schaefer and Whiticar [2008] would result in an ϵ_{tot} for GS-22 and GI-21.2 of -5.9‰ and -6.5‰ , respectively, if all other sink fluxes and ϵ values remained constant. This would increase the difference in $\delta^{13}\text{C}$ between the

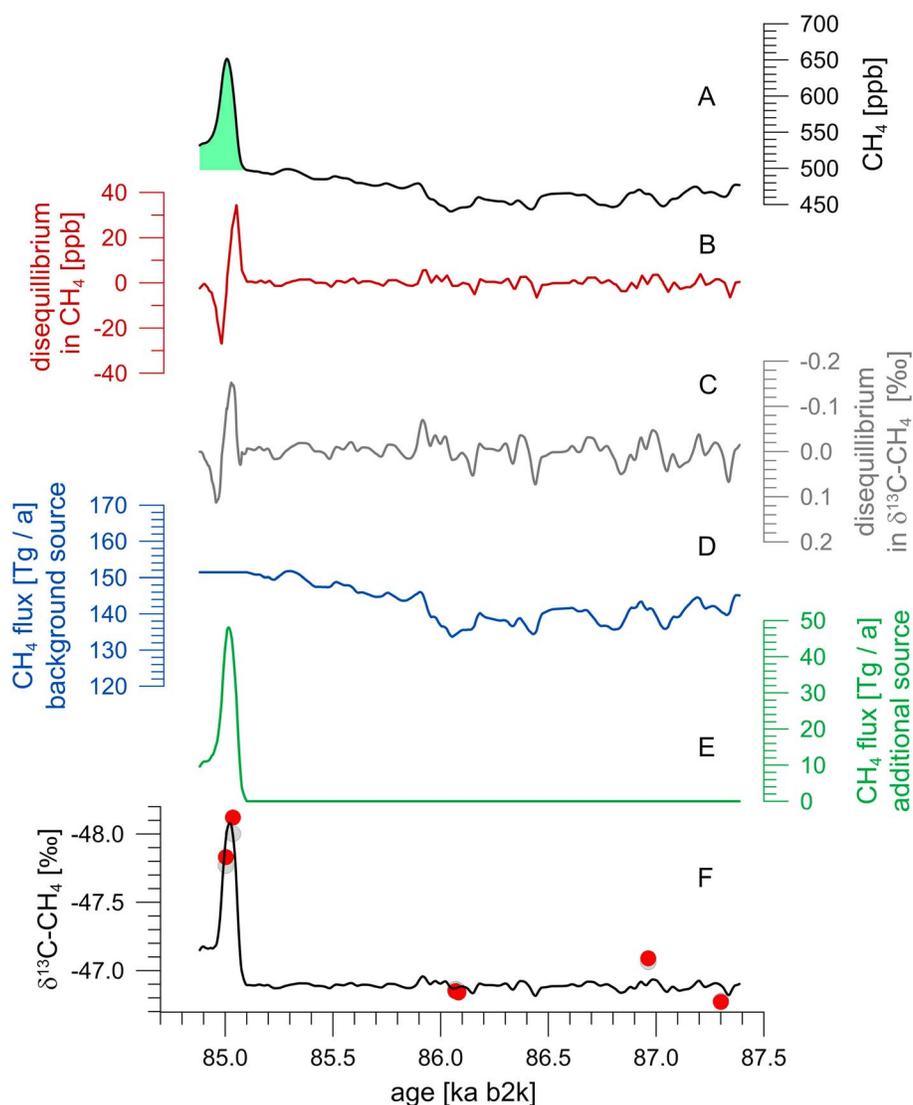


Figure 4. Box model data. (a) The $[\text{CH}_4]$ input data [Chappellaz *et al.*, 2013] with GI-21.2 highlighted in green. (b and c) The calculated disequilibrium effect for $[\text{CH}_4]$ and $\delta^{13}\text{C}-\text{CH}_4$, respectively. (d and e) The calculated CH_4 fluxes of the background source and the additional source during GI-21.2, respectively. (f) The $\delta^{13}\text{C}-\text{CH}_4$ output from the box model in comparison to our $\delta^{13}\text{C}-\text{CH}_4$ measurements before (red circles) and after (grey circles) the disequilibrium correction.

ϵ_{tot} -corrected GS-22 and GI-21.2 data by 0.6‰ and shift the $\delta^{13}\text{C}$ of the total sources toward more enriched values. The scaled ϵ_{tot} scenario would deplete the KPA results by 1.3‰. Note that none of these ϵ_{tot} scenarios would change our KPA results beyond the uncertainty envelope of $\pm 2\%$ and would therefore not change our interpretation of the results.

Applying the disequilibrium correction (equation (3)) shifts the KPA results by 0.4‰ toward stronger $\delta^{13}\text{C}$ enrichment, which is a small effect compared to the KPA uncertainty of $\pm 2\%$.

3.2. Comparing the Results From Keeling Plot Analysis and Box Model

The box model resulted in a $\delta^{13}\text{C}$ of the additional source that is 0.6‰ more depleted in $\delta^{13}\text{C}$ than the KPA result of the $[\text{CH}_4]$ -correlated $\delta^{13}\text{C}-\text{CH}_4$ background. For the $[\text{CO}_2]$ -correlated background scenario, the box model result is 0.4‰ more enriched in $\delta^{13}\text{C}$. The difference between KPA and box model results is of different sign for both scenarios but well within the $\pm 2\%$ uncertainty of the KPA. The disagreement between KPA and box model is partly due to the different methods used to determine the mathematical solution. While the KPA result is determined by a least squares fit that considers all data points from GI-21.2 and GS-22, the box model result depends on an equal mismatch between the modeled $\delta^{13}\text{C}-\text{CH}_4$ scenario and the two GI-21.2 data

Table 2. Average $\delta^{13}\text{C}$ Isotope Ratios of Categorized CH_4 Sources^a

| Source | $\delta^{13}\text{C}\text{-CH}_4$ (‰) |
|------------------------|---------------------------------------|
| Tropical wetlands | -58 ^b |
| Boreal wetlands | -63 ^b |
| CH_4 hydrates | -62.5 ^b |
| Aerobic C3 | -58 ^c |
| Aerobic C4 | -50 ^c |
| Termites | -70 ^d |
| Geological | -40 ^e |
| Biomass burning | -25 ^d |
| Thermokarst lakes | -70 ^f |

^aThe data represent average values with an uncertainty of 2–5‰ [e.g., Quay *et al.*, 1999] or even larger for CH_4 hydrates [e.g., Kvenvolden, 1995].

^bFrom Whiticar and Schaefer [2007] for glacial periods.

^cFrom Keppler *et al.* [2006].

^dFrom Mikaloff Fletcher *et al.* [2004].

^eFrom Denman *et al.* [2007].

^fFrom Walter *et al.* [2008].

points. Furthermore, the $[\text{CH}_4]$ at the beginning of the GI-21.2 event is ~ 20 ppb higher than the mean $[\text{CH}_4]$ of the background samples from GS-22. To compensate this larger proportion of $\delta^{13}\text{C}$ -enriched background CH_4 in the box model, the $\delta^{13}\text{C}$ of the additional CH_4 has to be more depleted. Note that this is not the case for the box model test with the $[\text{CO}_2]$ -correlated background, which may explain the change of sign in the difference.

3.3. Averaging the Keeling Plot Analyses Results of Both Background Assumptions for Interpretation

For our interpretation, we average the KPA results using the $[\text{CH}_4]$ - and the $[\text{CO}_2]$ -correlated background scenarios to $-56.8 \pm 2.8\%$. The averaged result considers both background scenarios as equally possible solutions within the propagated uncertainty range.

4. Discussion

4.1. Applicability of Keeling Plot Analysis to Study the Variation of $\delta^{13}\text{C}\text{-CH}_4$ in Ice Core Samples

We use KPA for the analysis of $\delta^{13}\text{C}\text{-CH}_4$ in ice core samples and derive results that agree with a forward stepping box model within the uncertainty of the KPA. For most accurate results, the KPA requires the disequilibrium correction, which can be calculated using a suitable box model in addition to the KPA. However, the disequilibrium correction did not change our KPA results significantly. The disequilibrium effect could be relatively small in our analysis because each of our $\delta^{13}\text{C}\text{-CH}_4$ samples averages a large part of the GI-21.2 event; hence, extreme values of the disequilibrium effect are smoothed out. This could be different for the analysis of samples with a similar but longer-lasting $[\text{CH}_4]$ gradient or samples that integrate a shorter period of time. However, the $[\text{CH}_4]$ growth rate during GI-21.2 is to date the highest observed for natural $[\text{CH}_4]$ variability [Chappellaz *et al.*, 2013]. Therefore, it seems unlikely that the disequilibrium effect will have a significant impact on a KPA that is performed on other ice core samples during other periods of time.

Besides the fact that KPA can serve as a fast approach for $\delta^{13}\text{C}\text{-CH}_4$ analysis, it has the advantage over a box model that it is independent of a gas age scale, as KPA only requires the combination of CH_4 and $\delta^{13}\text{C}\text{-CH}_4$ data. This avoids uncertainties from dating issues that may arise when data sets from more than one ice core are merged for analysis.

4.2. CH_4 Source Identification From Keeling Plot Analysis Result

Our averaged KPA result of $-56.8 \pm 2.8\%$ is in best agreement with the $\delta^{13}\text{C}$ that Whiticar and Schaefer [2007] reconstructed for CH_4 emissions from tropical wetlands during the glacial period (Table 2). Note that aerobic formation of CH_4 in C3 plants is a theoretical solution; however, the process and its relevance are not well understood and can therefore not be discussed in the context of rapid climate changes.

Our preferred solution is that tropical wetland emissions were the most important contributors to the $[\text{CH}_4]$ anomaly of GI-21.2. This result is supported by Baumgartner *et al.* [2014], who found that high growth rates, as reported for GI-21.2 [Chappellaz *et al.*, 2013], are associated with an Intertropical Convergence Zone (ITCZ) position that enhances CH_4 emissions from tropical Asia. Another method to constrain the spatial distribution of CH_4 emissions is the relative Inter-Polar Difference (rIPD) of CH_4 [e.g., Chappellaz *et al.*, 1997; Brook *et al.*, 2000; Baumgartner *et al.*, 2014], which is strongly influenced by solar insolation [e.g., Baumgartner *et al.*, 2014]. Because no rIPD reconstruction exists specifically for GI-21.2, we assume that the rIPD of the directly following GI-21.1e is indicative for GI-21.2. This assumption is justified by a maximum insolation gradient between 30°N and 60°N during both GI-21.1e and GI-21.2, with higher insolation in the lower latitudes [Laskar *et al.*, 2004].

The rIPD during GI-21.1e appears to be of medium amplitude with $\sim 7.5\%$, which indicates the dominance of tropical Northern Hemisphere CH_4 sources [Baumgartner *et al.*, 2014].

Predominant contributions from ^{13}C -depleted CH_4 sources like boreal wetlands would have to be compensated by emissions from ^{13}C -enriched CH_4 sources (such as pyrogenic CH_4) to match the isotope budget. Such a scenario cannot be ruled out by our data but is not consistent with the geographic constraints for the additional CH_4 emissions described above. The KPA precludes either biomass burning or termites as main CH_4 source for the GI-21.2 anomaly (Table 2). Interestingly, an outstanding CH_4 emission pulse from geological mantle sources can also be ruled out, as this CH_4 source is significantly more enriched in ^{13}C [e.g., *Etiopie and Lollar*, 2013]. The wide $\delta^{13}\text{C}$ range of marine hydrate-bound CH_4 between -57 and -73‰ [e.g., *Kvenvolden*, 1995] with the additional potential for postemission enrichment by microbial oxidation in the water column [e.g., *Whiticar and Faber*, 1986] does not rule out CH_4 hydrate destabilization as the cause of the $[\text{CH}_4]$ anomaly during GI-21.2.

To investigate this further, we calculate a Rayleigh distillation as described by *Schaefer et al.* [2006] and find a theoretical CH_4 hydrate emission scenario in which $\sim 47\%$ of released CH_4 is oxidized in sediment and water column while the remaining $\sim 53\%$ reach the atmosphere in agreement with our KPA result. However, this partitioning between CH_4 oxidation and release to the atmosphere is in strong disagreement with the observations of *Yvon-Lewis et al.* [2011], who show that only 0.01% of the CH_4 that was released during the “Deep Water Horizon” spill arrived in the atmosphere. Based on $\delta^2\text{H}-\text{CH}_4$ data, *Sowers* [2006] and *Bock et al.* [2010] proved that marine hydrate destabilization did not cause the $[\text{CH}_4]$ increase during the last deglaciation and of GI-7 and GI-8, respectively. For the above mentioned reasons, we assume that it is unlikely that CH_4 released from marine hydrates has caused the $[\text{CH}_4]$ variability during GI-21.2.

In general, it is important to remember that the complex biogeochemistry of natural CH_4 sources leads to wide $\delta^{13}\text{C}$ ranges within each source category, thereby limiting the constraining power of $\delta^{13}\text{C}-\text{CH}_4$ reconstructions. In the following, we compare our hypothesis that mostly tropical CH_4 sources have caused the $[\text{CH}_4]$ anomaly during GI-21.2 to independent evidence.

5. Comparison to Independent Climate Records

The strong temperature variability as recorded in Greenland ice cores during the last glacial (e.g., *NGRIP community members* [2004] and Figure 5c) is associated with changes in both atmospheric and ocean circulation [e.g., *Chiang and Friedman*, 2012]. Variations in both are of hemispheric extent and are related to the location of the ITCZ [*Chiang and Friedman*, 2012; *Burckel et al.*, 2015]. The ITCZ location is critical for the regulation of monsoon system intensities by controlling the meridional transport of heat and moisture [*Chiang and Friedman*, 2012; *Burckel et al.*, 2015], which are the main controlling measures of CH_4 emissions [*Guo et al.*, 2012]. Thus, the ITCZ location determines the meridional distribution of CH_4 source regions and thereby the partitioning of CH_4 emissions from the Northern and Southern Hemispheres [e.g., *Guo et al.*, 2012; *Baumgartner et al.*, 2014]. In the following, we discuss the variation of climate proxies in a geographical context in order to test our hypothesis that the CH_4 excursion during GI-21.2 was predominantly caused by increased CH_4 emissions from tropical wetlands.

5.1. South American Monsoon Systems During Greenland Interstadial 21.2

Wang et al. [2004], *Cruz et al.* [2005], and *Deplazes et al.* [2013] reconstruct large-scale precipitation variations in South American rainforest systems: During GS-22, the Atlantic rainforest section of southern South America experienced a wet interval [*Wang et al.*, 2004; *Cruz et al.*, 2005] (Figure 5e), thereby enhancing CH_4 emissions from this region in the Southern Hemisphere. At the same time, marine sediment reflectivity records from the Cariaco Basin indicate that the climate in the much larger Amazonian rainforest system was in a dryer phase [*Deplazes et al.*, 2013] (Figure 5d), which can be expected to reduce CH_4 emissions from the Amazonian region. After GS-22, the large-scale precipitation pattern over South America changed significantly, most likely driven by solar insolation and changes in ocean circulation [*Wang et al.*, 2004; *Burckel et al.*, 2015]. The transition between the two states is resolved in both the Cariaco Basin sediments [*Deplazes et al.*, 2013] and the subtropical speleothem records from south Brazil [*Cruz et al.*, 2005], where both records indicate a wetter climate period in the Amazonian rainforest systems during GI-21.2 (Figures 5d and 5e). *Deplazes et al.* [2013] propose a strong precipitation increase in the Amazonian rainforest that coincided with the temperature and $[\text{CH}_4]$ anomaly of GI-21.2, thereby linking Greenlandic and South American climate. The ITCZ controlled precipitation pattern in South America [*Wang et al.*, 2004; *Cruz et al.*, 2005; *Deplazes et al.*, 2013] may result in a change

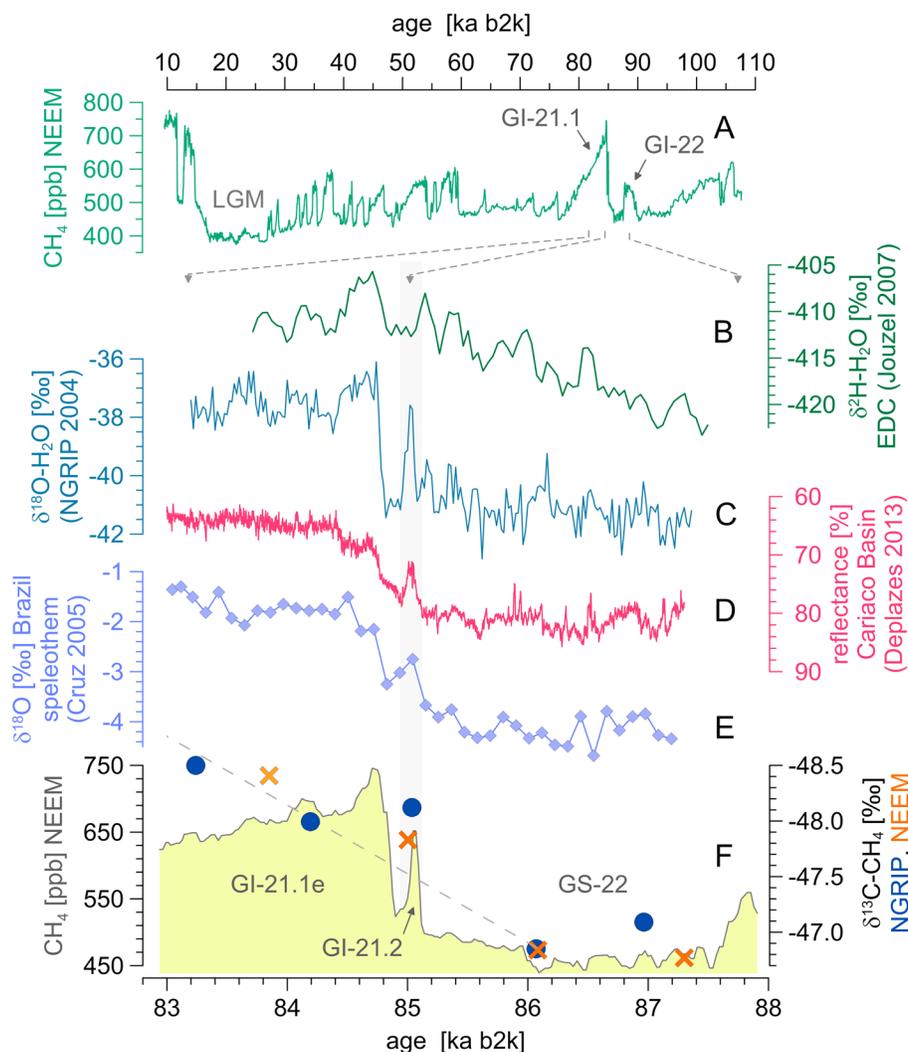


Figure 5. $\delta^{13}\text{C}\text{-CH}_4$ in comparison to independent climate data. (a) $[\text{CH}_4]$ during last glacial [Chappellaz *et al.*, 2013] on the top x axis. Figures 5b–5f refer to the bottom x axis. The grey bar highlights the GI-21.2 event. (b) Antarctic and (c) Greenlandic water isotope ratios [Jouzel *et al.* [2007] and NGRIP [2004], respectively). (d) Reflectivity record of Cariaco Basin [Deplazes *et al.*, 2013] and (e) $\delta^{18}\text{O}$ from south Brazilian speleothems [Cruz *et al.*, 2005]. (f) Left axis $[\text{CH}_4]$ and right axis (inverted) $\delta^{13}\text{C}\text{-CH}_4$ from NGRIP (circles) and NEEEM (crosses). All data are shown on the GICC05_modelext timescale [Wolff *et al.*, 2010]. $\delta^{18}\text{O}$ Figure 5e is adjusted by +0.55ka to align GI-21.2, which is in line with Cruz *et al.* [2005].

of the dominant South American CH_4 source regions from the smaller Atlantic to the much larger Amazonian region between GS-22 and GI-21.2, possibly triggering CH_4 fluxes from Amazonian wetland sources to cause the $[\text{CH}_4]$ anomaly of GI-21.2.

5.2. Asian Monsoon Systems During Greenland Interstadial 21.2

Pausata *et al.* [2011], Deplazes *et al.* [2013], and Mohtadi *et al.* [2014] propose a teleconnection pattern where North Atlantic cold periods coincided with dryer conditions in the tropical Indian monsoon realm, mediated by changes in atmospheric circulation. Speleothem records of Wang *et al.* [2001, 2008] provide geological evidence that link Greenland climate with the transport of heat and moisture in the distant East Asian monsoon region. A further geological argument for a tight coupling between Greenland temperature and the Asian monsoon system is suggested by Ruth *et al.* [2007], according to which the lower dust concentration observed in Greenland ice cores during GI-21.2 [e.g., Rasmussen *et al.*, 2014] indicates an intensification of the Asian monsoon system. The above mentioned studies suggest warmer and wetter climate in the Asian monsoon region during GI-21.2, where both factors potentially raise tropical wetland CH_4 emissions [e.g., Brook *et al.*, 1996; Guo *et al.*, 2012].

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5.3. Potential for Boreal and Pyrogenic CH₄ Emissions During Greenland Interstadial 21.2

Guo *et al.* [2012] describe that enhanced monsoon system strength (sections 5.1 and 5.2) increases the meridional transport of heat and moisture into the extratropics, thereby strengthening boreal CH₄ emissions. Also, high Northern Hemisphere summer insolation [Laskar *et al.*, 2004] and the relatively warm polar temperatures [EPICA, 2006] during GI-21.2 had the potential to enhance CH₄ emissions from boreal wetlands [e.g., Brook *et al.*, 1996]. However, the CH₄ of boreal wetland sources is relatively depleted in ¹³C and ranges around −63‰ (Table 2). Moreover, McCalley *et al.* [2014] report a further, systematic $\delta^{13}\text{C}$ depletion of CH₄ emissions from newly degrading permafrost soils, which we cannot rule out during the rapid Northern Hemisphere warming of GI-21.2 [NGRIP, 2004]. Therefore, our result of $-56.8 \pm 2.8\text{‰}$ precludes boreal wetland emissions as the predominant CH₄ source causing the [CH₄] anomaly of GI-21.2 (Table 2). Increased boreal CH₄ emissions would have to be compensated by ¹³C-enriched pyrogenic CH₄ emissions (−25‰) during the period integrated per $\delta^{13}\text{C-CH}_4$ sample to match our reconstructed $\delta^{13}\text{C-CH}_4$. Daniau *et al.* [2007] analyzed charcoal records and found increased wildfire intensity in Europe during interstadial periods of the last glacial period, which would be consistent with this scenario.

On the contrary, charcoal records from lower latitudes suggest minimum wildfire intensities during our study period [e.g., Bird and Cali, 1998; Wang *et al.*, 2005; Kershaw *et al.*, 2007]. Given that Thonicke *et al.* [2005] report a consistent latitudinal pattern of highest charcoal counts and pyrogenic CH₄ emissions in the lower latitudes and found that this pattern was more pronounced during the LGM, the lack of evidence for increased tropical burning during our study period suggests that global pyrogenic CH₄ formation was small. Consequently, CH₄ emissions from boreal wetland sources must have been small as well to meet the $\delta^{13}\text{C-CH}_4$ constraint. This finding strengthens our interpretation that tropical wetlands were the most important CH₄ sources that caused the [CH₄] anomaly of GI-21.2.

6. Conclusions

We present an approach to use Keeling plot analysis to investigate the variability of CH₄ sources in ice core samples, based on $\delta^{13}\text{C-CH}_4$ and [CH₄] measurements during GI-21.2. Our Keeling plot analysis includes a correction for the disequilibrium effect, based on information from a time stepping box model. The result of the Keeling plot analysis agrees with the box model results within 0.6‰, which is well within the uncertainty of the Keeling plot analysis ($\pm 2\text{‰}$) for either of the background scenarios.

The average $\delta^{13}\text{C-CH}_4$ of the source causing the rapid [CH₄] variability is best matched by enhanced CH₄ emissions from tropical wetlands. Our conclusion is supported by a range of independent climate records, which suggest wetter and warmer climate in tropical Amazonian and Asian CH₄ source ecosystems. Increases in both boreal wetlands and biomass burning were only possible if their relative contributions produced the $\delta^{13}\text{C}$ of CH₄ sources suggested by the KPA. Because charcoal records from lower latitudes suggest that the wildfire intensity during our study period was low, a strong response of boreal CH₄ sources to the rapid GI-21.2 event seems unlikely. The hypothesis that boreal CH₄ sources showed a low sensitivity to the short but rapid temperature increase of GI-21.2 [e.g., NGRIP, 2004; Capron *et al.*, 2010; Boch *et al.*, 2011] is an interesting finding in the light of global warming and the associated potential for CH₄ emissions from boreal sources such as permafrost (note that GI-21.2 is not a complete analogue for present-day climate conditions). Further research including climate-vegetation modeling and atmospheric observations at highest spatiotemporal resolution is needed to provide reliable scenarios of vegetation dynamics and related changes in isotope ratios of CH₄ emissions in order to fully exploit the information provided by CH₄ isotope records from ice cores.

References

- Baumgartner, M., *et al.* (2014), NGRIP CH₄ concentration from 120 to 10 kyr before present and its relation to a $\delta^{15}\text{N}$ temperature reconstruction from the same ice core, *Clim. Past*, 10, 903–920.
- Bereiter, B., D. Lüthi, M. Siegrist, S. Schüpbach, T. F. Stocker, and H. Fischer (2012), Mode change of millennial CO₂ variability during the last glacial cycle associated with a bipolar marine carbon seesaw, *Proc. Natl. Acad. Sci. U.S.A.*, 109(25), 9755–9760.
- Bird, M. I., and J. A. Cali (1998), A million-year record of fire in sub-Saharan Africa, *Nature*, 394, 767–769.
- Blunier, T., R. Spahni, J. M. Barnola, J. Chappellaz, L. Loulergue, and J. Schwander (2007), Synchronization of ice core records via atmospheric gases, *Clim. Past*, 3(2), 325–330.
- Boch, R., H. Cheng, C. Spotl, R. L. Edwards, X. Wang, and P. Hauselmann (2011), NALPS: A precisely dated European climate record 120–60 ka, *Clim. Past*, 7(4), 1247–1259.
- Bock, M., J. Schmitt, L. Möller, R. Spahni, T. Blunier, and H. Fischer (2010), Hydrogen isotopes preclude marine hydrate CH₄ emissions at the onset of Dansgaard-Oeschger events, *Science*, 328(5986), 1686–1689.

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- Brook, E. J., T. Sowers, and J. Orchardo (1996), Rapid variations in atmospheric methane concentration during the past 110,000 years, *Science*, 273(5278), 1087–1091.
- Brook, E. J., S. Harder, J. Severinghaus, E. J. Steig, and C. M. Sucher (2000), On the origin and timing of rapid changes in atmospheric methane during the last glacial period, *Global Biogeochem. Cycles*, 14(2), 559–572.
- Buizert, C., et al. (2012), Gas transport in firn: Multiple-tracer characterisation and model intercomparison for NEEM, Northern Greenland, *Atmos. Chem. Phys.*, 12(9), 4259–4277.
- Buizert, C., T. Sowers, and T. Blunier (2013), Assessment of diffusive isotopic fractionation in polar firn, and application to ice core trace gas records, *Earth Planet. Sci. Lett.*, 361, 110–119.
- Burckel, P., C. Waelbroeck, J. M. Gherardi, S. Pichat, H. Arz, J. Lippold, T. Dokken, and F. Thil (2015), Atlantic Ocean circulation changes preceded millennial tropical South America rainfall events during the last glacial, *Geophys. Res. Lett.*, 42, 411–418.
- Capron, E., et al. (2010), Millennial and sub-millennial scale climatic variations recorded in polar ice cores over the last glacial period, *Clim. Past*, 6(3), 345–365.
- Chappellaz, J., T. Blunier, S. Kints, A. Dällenbach, J. M. Barnola, J. Schwander, D. Raynaud, and B. Stauffer (1997), Changes in the atmospheric CH₄ gradient between Greenland and Antarctica during the Holocene, *J. Geophys. Res. Atmos.*, 102(D13), 15987–15997.
- Chappellaz, J., et al. (2013), High-resolution glacial and deglacial record of atmospheric methane by continuous-flow and laser spectrometer analysis along the NEEM ice core, *Clim. Past*, 9, 2579–2593.
- Chiang, J. C., and A. R. Friedman (2012), Extratropical cooling, interhemispheric thermal gradients, and tropical climate change, *Annu. Rev. Earth Pl. Sc.*, 40, 383–412.
- Cruz, F. W., S. J. Burns, I. Karmann, W. D. Sharp, M. Vuille, A. O. Cardoso, J. A. Ferrari, P. L. S. Dias, and O. Viana (2005), Insolation-driven changes in atmospheric circulation over the past 116,000 years in subtropical Brazil, *Nature*, 434(7029), 63–66.
- Daniau, A. L., M. F. Sanchez-Goni, L. Beaufort, F. Laggoun-Defarge, M. F. Loutre, and J. Duprat (2007), Dansgaard-Oeschger climatic variability revealed by fire emissions in southwestern Iberia, *Quat. Sci. Rev.*, 26, 1369–1383.
- Denman, K. L., et al. (2007), Couplings between changes in the climate system and biogeochemistry, in *Climate Change 2007: The Physical Science Basis. Contribution of Working Group I to the Fourth Assessment Report of the Intergovernmental Panel on Climate Change*, edited by S. Solomon et al., pp. 499–587, Cambridge Univ. Press, Cambridge, U. K., and New York.
- Deplazes, G., et al. (2013), Links between tropical rainfall and North Atlantic climate during the last glacial period, *Nat. Geosci.*, 6(3), 213–217.
- EPICA community members (2006), One-to-one coupling of glacial climate variability in Greenland and Antarctica, *Nature*, 444(7116), 195–198.
- Etheridge, D. M., L. P. Steele, R. J. Francey, and R. L. Langenfelds (1998), Atmospheric methane between 1000 A.D. and present: Evidence of anthropogenic emissions and climatic variability, *J. Geophys. Res.*, 103(D13), 15,979–15,993.
- Etiope, G., and B. S. Lollar (2013), Abiotic methane on Earth, *Rev. Geophys.*, 51(2), 276–299.
- Ferretti, D. F., et al. (2005), Unexpected changes to the global methane budget over the past 2000 years, *Science*, 309(5741), 1714–1717.
- Fischer, H., et al. (2008), Changing boreal methane sources and constant biomass burning during the last termination, *Nature*, 452(7189), 864–867.
- Fisher, R. E., et al. (2011), Arctic methane sources: Isotopic evidence for atmospheric inputs, *Geophys. Res. Lett.*, 38, L21803, doi:10.1029/2011GL049319.
- Forster, P., et al. (2007), Changes in atmospheric constituents and in radiative forcing, in *Climate Change 2007: The Physical Science Basis. Contribution of Working Group I to the Fourth Assessment Report of the Intergovernmental Panel on Climate Change*, edited by S. Solomon et al., pp. 129–234, Cambridge Univ. Press, Cambridge, U. K., and New York.
- Francey, R. J., M. R. Manning, C. E. Allison, S. A. Coram, D. M. Etheridge, R. L. Langenfelds, D. C. Lowe, and L. P. Steele (1999), A history of $\delta^{13}\text{C}$ in atmospheric CH₄ from the Cape Grim Air Archive and Antarctic firn air, *J. Geophys. Res. Atmos.*, 104(D19), 23,631–23,643.
- Grachev, A. M., E. J. Brook, and J. P. Severinghaus (2007), Abrupt changes in atmospheric methane at the MIS 5b-5a transition, *Geophys. Res. Lett.*, 34, L20703, doi:10.1029/2007GL029799.
- Guo, Z. T., X. Zhou, and H. B. Wu (2012), Glacial-interglacial water cycle, global monsoon and atmospheric methane changes, *Clim. Dynam.*, 39(5), 1073–1092.
- Jouzel, J., et al. (2007), Orbital and millennial Antarctic climate variability over the past 800,000 years, *Science*, 317(5839), 793–796.
- Keeling, C. D. (1958), The concentration and isotopic abundances of atmospheric carbon dioxide in rural areas, *Geochim. Cosmochim. Ac.*, 13, 322–334.
- Keppler, F., J. T. G. Hamilton, M. Brass, and T. Röckmann (2006), Methane emissions from terrestrial plants under aerobic conditions, *Nature*, 439(7073), 187–191.
- Kershaw, P. A., S. C. Bretherton, and S. van der Kaars (2007), A complete pollen record of the last 230 ka from Lynch's Crater, north-eastern Australia, *Palaeogeogr. Palaeoclimatol.*, 251, 23–45.
- Kvenvolden, K. A. (1995), A review of the geochemistry of methane in natural gas hydrate, *Org. Geochem.*, 23, 997–1008.
- Laskar, J., P. Robutel, F. Joutel, M. Gastineau, A. C. M. Correia, and B. Levrard (2004), A long-term numerical solution for the insolation quantities of the Earth, *Astron. Astrophys.*, 428, 261–285.
- Lassey, K. R., D. M. Etheridge, D. C. Lowe, A. M. Smith, and D. F. Ferretti (2007), Centennial evolution of the atmospheric methane budget: What do the carbon isotopes tell us?, *Atmos. Chem. Phys.*, 7, 2119–2139.
- Levine, J. G., E. W. Wolff, P. O. Hopcroft, and P. J. Valdes (2012), Controls on the tropospheric oxidizing capacity during an idealized Dansgaard-Oeschger event, and their implications for the rapid rises in atmospheric methane during the last glacial period, *Geophys. Res. Lett.*, 39, L12805, doi:10.1029/2012GL051866.
- Loulergue, L., A. Schilt, R. Spahni, V. Masson-Delmotte, T. Blunier, B. Lemieux, J. M. Barnola, D. Raynaud, T. F. Stocker, and J. Chappellaz (2008), Orbital and millennial-scale features of atmospheric CH₄ over the past 800,000 years, *Nature*, 453(7193), 383–386.
- McCalley, C. K., B. J. Woodcroft, S. B. Hodgkins, R. A. Wehr, E.-H. Kim, R. Mondav, P. M. Crill, J. P. Chanton, V. I. Rich, G. W. Tyson, and S. R. Saleska (2014), Methane dynamics regulated by microbial community response to permafrost thaw, *Nature*, 453(7193), 383–386.
- Meehl, G. A., et al. (2007), Couplings between changes in the climate system and biogeochemistry, in *Climate Change 2007: The Physical Science Basis. Contribution of Working Group I to the Fourth Assessment Report of the Intergovernmental Panel on Climate Change*, edited by S. Solomon et al., pp. 499–587, Cambridge Univ. Press, Cambridge, U. K., and New York.
- Melton, J. R., H. Schaefer, and M. J. Whiticar (2012), Enrichment in $\delta^{13}\text{C}$ of atmospheric CH₄ during the Younger Dryas termination, *Clim. Past*, 8, 1177–1197.
- Mikaloff Fletcher, S. E., P. P. Tans, L. M. Bruhwiler, J. B. Miller, and M. Heimann (2004), CH₄ sources estimated from atmospheric observations of CH₄ and its $^{13}\text{C}/^{12}\text{C}$ isotopic ratios: 1. Inverse modeling of source processes, *Global Biogeochem. Cycles*, 18, GB4004, doi:10.1029/2004GB002223.

- Mischler, J. A., T. A. Sowers, R. B. Alley, R. B. M. Battle, J. R. McConnell, L. Mitchell, T. Popp, E. Sofen, and M. K. Spencer (2009), Carbon and hydrogen isotopic composition of methane over the last 1000 years, *Global Biogeochem. Cycles*, 23, Gb4024, doi:10.1029/2009GB003460.
- Mitchell, L., E. Brook, J. E. Lee, C. Buizert, and T. Sowers (2013), Constraints on the Late Holocene anthropogenic contribution to the atmospheric methane budget, *Science*, 342(6161), 964–966.
- Mohtadi, M., M. Prange, D. W. Oppo, R. De Pol-Holz, U. Merkel, X. Zhang, S. Steinke, and A. Lückge (2014), North Atlantic forcing of tropical Indian Ocean climate, *Nature*, 509, 76–80.
- Möller, L., T. Sowers, M. Bock, R. Spahni, M. Behrens, J. Schmitt, H. Miller, and H. Fischer (2013), Independent variations of CH₄ emissions and isotopic composition over the past 160,000 years, *Nat. Geosci.*, 6(10), 885–890.
- NGRIP community members (2004), High-resolution record of Northern Hemisphere climate extending into the last interglacial period, *Nature*, 431(7005), 147–151.
- Pataki, D. E., J. R. Ehleringer, L. B. Flanagan, D. Yakir, D. R. Bowling, C. J. Still, N. Buchmann, J. O. Kaplan, and J. A. Berry (2003), The application and interpretation of Keeling plots in terrestrial carbon cycle research, *Global Biogeochem. Cycles*, 17(1), 1022, doi:10.1029/2001GB001850.
- Pausata, F. S. R., D. S. Battisti, K. H. Nisancioglu, and C. M. Bitz (2011), Chinese stalagmite $\delta^{18}\text{O}$ controlled by changes in the Indian monsoon during a simulated Heinrich event, *Nat. Geosci.*, 4(7), 474–480.
- Quay, P., J. Stutsman, D. Wilbur, A. Snover, E. Dlugokencky, and T. Brown (1999), The isotopic composition of atmospheric methane, *Global Biogeochem. Cycles*, 13(2), 445–461.
- Rasmussen, S. O., et al. (2014), A stratigraphic framework for abrupt climatic changes during the last glacial period based on three synchronized Greenland ice-core records: Refining and extending the INTIMATE event stratigraphy, *Quat. Sci. Rev.*, 106, 14–28.
- Ruth, U., M. Bigler, R. Röthlisberger, M. L. Siggaard-Andersen, S. Kipfstuhl, K. Goto-Azuma, M. E. Hansson, S. J. Johnsen, H. Y. Lu, and J. P. Steffensen (2007), Ice core evidence for a very tight link between North Atlantic and east Asian glacial climate, *Geophys. Res. Lett.*, 34, L03706, doi:10.1029/2006GL027876.
- Sapart, C. J., et al. (2011), Simultaneous stable isotope analysis of methane and nitrous oxide on ice core samples, *Atmos. Meas. Tech.*, 4(12), 2607–2618.
- Sapart, C. J., et al. (2012), Natural and anthropogenic variations in methane sources during the past two millennia, *Nature*, 490(7418), 85–88.
- Sapart, C. J., et al. (2013), Can the carbon isotopic composition of methane be reconstructed from multi-site firm air measurements?, *Atmos. Chem. Phys.*, 13(14), 6993–7005.
- Schaefer, H., and M. J. Whiticar (2008), Potential glacial-interglacial changes in stable carbon isotope ratios of methane sources and sink fractionation, *Global Biogeochem. Cycles*, 22, GB1001, doi:10.1029/2006GB002889.
- Schaefer, H., M. J. Whiticar, E. J. Brook, V. V. Petrenko, D. F. Ferretti, and J. P. Severinghaus (2006), Ice record of $\delta^{13}\text{C}$ for atmospheric CH₄ across the Younger Dryas-Preboreal transition, *Science*, 313(5790), 1109–1112.
- Schilt, A., et al. (2010), Atmospheric nitrous oxide during the last 140,000 years, *Earth Planet. Sci. Lett.*, 300, 33–43.
- Schmitt, J., B. Seth, M. Bock, H. Fischer, L. Möller, C. J. Sapart, M. Prokopiou, C. Van der Veen, T. Röckmann, and T. Sowers (2013), On the interference of Kr during carbon isotope analysis of atmospheric methane using continuous flow combustion—Isotope ratio mass spectrometry, *Atmos. Meas. Tech.*, 6, 1425–1445.
- Sowers, T. (2006), Late quaternary atmospheric CH₄ isotope record suggests marine clathrates are stable, *Science*, 311(5762), 838–840.
- Spahni, R., J. Schwander, J. Flückiger, B. Stauffer, J. Chappellaz, and D. Raynaud (2003), The Attenuation of Fast Atmospheric CH₄ Variations Recorded in Polar Ice Cores, *Geophys. Res. Lett.*, 30(11), 1571, doi:10.1029/2003GL017093.
- Sperlich, P., M. Guillevic, C. Buizert, T. M. Jenk, C. J. Sapart, H. Schaefer, T. J. Popp, and T. Blunier (2012), A combustion setup to precisely reference $\delta^{13}\text{C}$ and $\delta^2\text{H}$ isotope ratios of pure CH₄ to produce isotope reference gases of $\delta^{13}\text{C}$ -CH₄ in synthetic air, *Atmos. Meas. Tech.*, 5(9), 2227–2236.
- Sperlich, P., C. Buizert, T. M. Jenk, C. J. Sapart, M. Prokopiou, T. Röckmann, and T. Blunier (2013), An automated GC-C-GC-IRMS setup to measure paleoatmospheric $\delta^{13}\text{C}$ -CH₄, $\delta^{15}\text{N}$ -N₂ and $\delta^{18}\text{O}$ -N₂O in one ice core sample, *Atmos. Meas. Tech.*, 6(8), 2027–2041.
- Stowasser, C., et al. (2012), Continuous measurements of methane mixing ratios from ice cores, *Atmos. Meas. Tech.*, 5(5), 999–1013.
- Tans, P. P. (1997), A note on isotopic ratios and the global atmospheric methane budget, *Global Biogeochem. Cycles*, 11(1), 77–81.
- Thonicke, K., C. I. Prentice, and C. Hewitt (2005), Modeling glacial-interglacial changes in global fire regimes and trace gas emissions, *Global Biogeochem. Cycles*, 19, GB3008, doi:10.1029/2004GB002278.
- Vallelonga, P., et al. (2012), Duration of Greenland Stadial 22 and ice-gas Delta age from counting of annual layers in Greenland NGRIP ice core, *Clim. Past*, 8(6), 1839–1847.
- Walter, K. M., J. P. Chanton, F. S. Chapin, E. A. G. Schuur, and S. A. Zimov (2008), Methane production and bubble emissions from Arctic lakes: Isotopic implications for source pathways and ages, *J. Geophys. Res. Biogeo.*, 113, G00a08, doi:10.1029/2007JG000569.
- Wang, X., P. A. Peng, and Z. L. Ding (2005), Black carbon records in Chinese Loess Plateau over the last two glacial cycles and implications for paleofires, *Palaeogeogr. Palaeoclimatol.*, 223, 9–19.
- Wang, X. F., A. S. Auler, R. L. Edwards, H. Cheng, P. S. Cristalli, P. L. Smart, D. A. Richards, and C. C. Shen (2004), Wet periods in northeastern Brazil over the past 210 kyr linked to distant climate anomalies, *Nature*, 432(7018), 740–743.
- Wang, Y. J., H. Cheng, R. L. Edwards, Z. S. An, J. Y. Wu, C. C. Shen, and J. A. Dorale (2001), A high-resolution absolute-dated Late Pleistocene monsoon record from Hulu Cave, China, *Science*, 294(5550), 2345–2348.
- Wang, Y. J., H. Cheng, R. L. Edwards, X. G. Kong, X. H. Shao, S. T. Chen, J. Y. Wu, X. Y. Jiang, X. F. Wang, and Z. S. An (2008), Millennial- and orbital-scale changes in the East Asian monsoon over the past 224,000 years, *Nature*, 451(7182), 1090–1093.
- Whiticar, M., and E. Faber (1986), Methane oxidation in sediment and water column environments—Isotopic evidence, *Org. Geochem.*, 10, 759–768.
- Whiticar, M., and H. Schaefer (2007), Constraining past global tropospheric methane budgets with carbon and hydrogen isotope ratios in ice, *Philos. Trans. R. Soc. A*, 365(1856), 1793–1828.
- Wolff, E. W., J. Chappellaz, T. Blunier, S. O. Rasmussen, and A. Svensson (2010), Millennial-scale variability during the last glacial: The ice core record, *Quat. Sci. Rev.*, 29, 2828–2838.
- Yvon-Lewis, S. A., L. Hu, and J. Kessler (2011), Methane flux to the atmosphere from the Deepwater Horizon oil disaster, *Geophys. Res. Lett.*, 38, 10.1029/2010GL045928.