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Ed Brooks

c/o Climate of the Past

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Revision of manuscript cp-2019-109

Dear Ed,

unfortunately the revision of our manuscript has taken substantially longer than we had anticipated. Contrary to our expectations, we could not use the offline land surface model to do the sensitivity experiments, but had to redo all sensitivity experiments using the full Earth System Model.

I apologize for the delay – and am very glad I am finally able to send the revised manuscript to you.

As you will be able to see, the main change to the manuscript is that we performed additional sensitivity experiments to better explain the reasons for the change in methane emissions between LGM and PI. We analyse the results in an additional appendix (now appendix A, the old one became appendix B) and added some text to the experiment description section, as well as to the section on the wetland CH₄ emissions.

Of course there are further smaller changes in response to the reviewers' comments, which I will detail below

To this letter I have appended a detailed response to the reviewers' comments and a manuscript version that shows all changes. Please do let me know if you believe further changes to the manuscript will be necessary, or if you need any additional clarification.

Thank you very much.

All the best, Thomas Kleinen

PS: I will contact you separately by email about the James Lee et al. manuscript – I didn't think it necessary for this manuscript, but we are working on another one where we will also discuss the interpolar gradient in more detail, and there I will definitely want more information.

Reply to Reviewers

We very much thank the reviewers for taking the time to review our manuscript. I have included the reviewer's comments in bold font, while our our original reply is in normal font. Finally, what we have changed in the manuscript in order to address the reviewer's point is explained in blue colour.

Reviewer #1

The authors present a comprehensive suite of ESM simulations including methane emissions. They focus in 5 ka intervals from the LGM to the present day. They show that emissions are reduced by around 50% at the LGM relative to the PI, meaning that no atmospheric lifetime change is required to explain the observed low concentration at the LGM, consistent with 3 recent studies of the atmospheric chemistry of the LGM. This is the first study to reconcile the emissions required by the observed CH4 drop without requiring to a large change in lifetime. Hence, these results will be important for understanding CH4 during ice-ages. I recommend minor revisions as explained below.

Main comments

To me, the main results of the paper require more explanation. If the model is able to correctly resolve a 50% emissions reduction at the LGM relative to the PI, there may be different reasons for this, e.g.: (i) the simulated LGM climate is less favourable for emissions than in older studies; (ii) the CH4 model is more sensitive to LGM conditions than in older studies, either through a different tuning, or because of increased model realism/complexity. (iii) the coupling of the methane model to the ESM increases the sensitivity to LGM conditions. In my opinion these options need to be clearly evaluated, otherwise we only know that it is possible to get this 50% reduction but not why or how.

Yes, we agree that this is the major shortcoming of our manuscript. While we will not be able to explain why older studies did not get similar results – we do not have access to their models after all – we should be able to shed some more light on the reasons why our model does what it does. This will require further model experiments however, in order to isolate the different factors. At the time we originally wrote the manuscript, we did not have a working offline configuration of the land surface model available, so we couldn't perform the necessary sensitivity experiments. In the mean time, we have been able to solve the issues with the offline model, allowing us to run the necessary sensitivity experiments.

We have addressed this point by reviewer #1 by performing additional sensitivity sensitivity experiments. As it turned out that soils in the offline model are substantially dryer then in the online model, we could not use the offline land surface scheme, as originally thought, but had to perofrm additional experiments with the full ESM. Results of the experiments are analysed in appendix A, and additional text explaining the experiments and reporting the main results was added to sections 2.5 and 3.3.

Moreover, these results don't show increasing southern hemisphere emissions over the past 6ka that would explain the observed recovery of atmospheric CH4, thus meaning that a human influence is not required. It would be good to understand how and why your results don't replicate this result from Singarayer et al., 2011. You discuss this a bit in lines 397-402, but I believe this could benefit from more detail, without needing further simulations.

We will clarify this in the revised version.

Not having access to Singrayer's original results make a detailed comparison impossible. However we have added some discussion of this in section 3.7. We also see a decrease in SH emissions for 5 ka BP, but the change in total emissions for the time slice is dominated by the increase in North African emissions due to the monsoon increase, which likely is too strong in our model, although we are glad to finally get a stronger monsoon (including a green Sahara), something most other models do not reproduce.

Technical comments

Line 1: Sorry to be picky, but past tense would make more sense: "Underwent", "nearly doubled" etc.

Thanks for being picky – we will address this. Changed in text.

Line 50: "However, Levine et al. (2011) found very small changes in CH4 lifetime between LGM and PI"

You could mention here that this has also been found by Murray et al 2014, and Hopcroft et al 2017, using different atmosphere-chemistry models. It would also be worth pointing out here why the lifetime didn't change. i.e. because of balance of competing influences from changes in reactive compounds like isoprene and reduced reaction rates due to lower temperatures, and lower generation of OH from lower water vapour levels.

Thanks, we will include that in the revision. Added to text.

Line 54: Here you can set up the main finding of you paper, by pointing out here that none of these earlier studies have managed to get the right change in emissions and lifetime together. Very good point by the reviewer, we did exactly that.

Line 177: insured -> ensured Changed in text.

Line 313: delete 'in' Changed in text.

Line 397: This is really interesting, but why/how does your model manage this and previous studies did not? I realise this is complex, but are there any obvious difference in your simulated LGM state? Is it drier in critical areas? Or is it due to the added complexity of your wetland module?

As detailed above, we will perform some additional offline sensitivity experiments to address this. My assumption is that it's a combination of the relatively large change in soil carbon stocks and the temperature sensitivity of methane production that we included, which is relatively new and likely wasn't included in at least some of the previous modeling studies.

Addressed by additional sensitivity experiments.

Line 399: This is an interesting finding. What is the explanation though? Do you see the same reduction in precipitation and hence wetland flux in Southern hemisphere as found by Singarayer et al 2011?

We will analyse this in more detail and discuss it in the revised paper. See above – briefly discussed in section 3.7.

Line 433: I think I understand the meaning of this sentence: "In total the wetland emissions account for 93–96% of the net CH4 flux, and all other methane sources are of minor importance.", but perhaps you could reword it.

Thank you for pointing this out, we will clarify it. We have reformulated the sentence – I hope it is clearer now.

Figures: It is slightly surprising to see Termite CH4 emissions in Europe.

We agree. However it is what we get when we implement the termite model from the Kirschke et al. / Saunois et al. reviews. We may add a short discussion of this to the section. In the end we did not add any text on this, as Saunois et al. also show emissions in Europe in their paper — our model reproduces their distribution, so we didn't think any additional comment was required.

References L. Murray, L.J. Mickley, J.O. Kaplan, E.D. Sofen, M. Pfeiffer and B. Alexan-der (2014). Factors controlling variability in the oxidative capacity of the troposphere since the Last Glacial Maximum, Atmos. Chem. Phys. 14, 3589-3622.

Reviewer #2

Manuscript summary: Thomas Kleinen et al. present an analysis of changes in methane fluxes from wetlands, termites and wildfires since the LGM. The analysis is performed using the Max Planck Institute for Meteorology Earth System Model, which explicitly simulates methane emissions (and the soil sink). Time-slice experiments are performed in the model, at 5 kyr intervals beginning at 20 kyr. The model is also run for the present day and compared with best available methane budget assessments. The authors find that wetland methane emissions dominated the changes in atmospheric methane over this time, and that tropical wetlands were the most important component of this.

Overall assessment and major comments:

It is difficult for me to assess the technical aspects of the MPI-ESM work, as I do not work with ESMs myself; I hope that another reviewer is able to do this. That said, the provided descriptions suggest a comprehensive and well-grounded approach, and the MPI Meteorology group does very good work in my opinion. The model simulates present-day methane emissions that are reasonable and generally compare well with top-down and bottom-up constraints. The model also produces methane emissions that appear to be mostly consistent with the ice core atmospheric methane record. My main concern with this submission to CP is its relative lack of novelty. I view CP as one of the leading journals publishing on paleoclimate, and as such I think that successful submissions to this journal should add substantially to our understanding of some aspect of paleoclimate. The major finding of the paper (that tropical wetland emissions were the main

factor driving the LGM - PI atmospheric methane change) has been argued for many times previously, including by model-based studies. While there have been studies arguing for other factors (e.g., the Kaplan et al 2006 study the authors cited), the leading role of tropical wetlands is the most accepted explanation. I think additional model results are valuable, even if they only reinforce the currently accepted hypothesis, but I'm not sure that CP is the best place – Earth System Science Data may be a better fit for this kind of study.

It may be possible that the work described in this manuscript is much more technically advanced than prior efforts. In this case, a publication in CP may be warranted, but the authors should then make a very clear argument for why their model is superior to what has been done before, and is expected to produce the most reliable results. Additional comments: I would recommend the addition of ice core constraints regarding the methane interpolar gradient (e.g., Baumgartner et al., 2012, Biogeosciences) into the analysis – is the partitioning between tropical and extratropical sources in the model consistent with these constraints?

We very much thank the reviewer for the overall praise that we read from her or his comments. However we have to disagree in some aspects:

Yes, the reviewer is perfectly correct that our finding that tropical wetlands are the dominant source of methane is not novel in itself. However, to our knowledge nobody has been able to show this in Earth System Model results, certainly not in a setup as internally consistent as ours.

We are able to show that we get reasonable emissions for the present-day situation, including a latitudinal distribution that is consistent with atmospheric inversions. Most other studies that we are aware of were not able to show this. We are also able to show that our emissions for other time slices are reasonable, in the sense that they are similar enough to ice core reconstructions to fall within a quantified uncertainty range, and we do not require major adjustments of the atmospheric lifetime of methane in order to achieve this.

We therefore argue that our results are more technically advanced than previous efforts. We will also add a more thorough analysis on the reasons why we do get these better results than previous studies, as detailed in the reply to referee #1.

The reviewer's point about the interpolar gradient, however, we regard as a very good suggestion, we will certainly take it up in the revision.

We addressed the point about the novelty and relevance of our results by adding a sentence in the introduction section, "none of the previous studies have managed to obtain the required changes in methane emissions, while fulfilling the constraints of the present-day methane budget at the same time."

We have also added a discussion of the implications of the interpolar gradient in section 3.7, citing Baumgartner and Mitchell.

Page 7, last paragraph (around line 210). The disagreement between model results and satellite observations for surface inundation is discouraging. I would recommend more discussion regarding how much uncertainty / error this could potentially introduce into the model wetland emissions estimates.

We will attempt to do so. Unfortunately the current state of remote sensing of inundation under closed canopies, as in tropical rainforests, leaves much to be desired, and major discrepancies exist between different remote sensing products. We acknowledge that we did discuss this well, we will attempt to improve it for the revised version.

We have attempted to improve this, but are somewhat unhappy with the result – we think we clarified this a little, but not comprehensively, but doing better would have required a lengthy discussion of

remote sensing products, their uncertainties and their interpretation, and we felt the issue does not warrant a lengthy discussion as it isn't the focus of the present paper.

Minor comments: Line 15-17. The Oldest Dryas – Bolling was an interval of similarly rapid methane change, I recommend mentioning this

Thank you for reminding us, we will do so. Added to the introduction section.

Paragraph around line 50. I would recommend adding the GESO-Chem LGM and PI results of Murray et al., 2014, ACP into the discussion of methane lifetime.

Yes, thank you for reminding us. Somehow the Murray et al. reference was lost in one of the previous revisions of the manuscript, we will add it again. Added to the introduction section.

Terrestrial methane emissions from Last Glacial Maximum to preindustrial

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Abstract. We investigate the changes in terrestrial natural methane emissions between the Last Glacial Maximum (LGM) and preindustrial (PI) by performing time-slice experiments with a methane-enabled version of MPI-ESM, the Max Planck Institute for Meteorology Earth System Model. We consider all natural sources of methane except for emissions from wild animals and geological sources, i.e. emissions from wetlands, fires, and termites. Changes are dominated by changes in tropical wetland emissions, with mid-to-high latitude wetlands playing a secondary role, and all other natural sources being of minor importance. The emissions are determined by the interplay of vegetation productivity, a function of CO₂ and temperature, source area size, affected by sea level and ice sheet extent, and the state of the West African Monsoon, with increased emissions from north Africa during strong monsoon phases.

We show that it is possible to explain the difference in atmospheric methane between LGM and PI purely by changes in emissions. As emissions more than double between LGM and PI, changes in the atmospheric lifetime of CH₄, as proposed in other studies, are not required.

1 Introduction

The atmospheric concentration of methane undergoes underwent major changes in the time between the last glacial maximum (LGM) and preindustrial (PI). Between LGM and 10 ka BP (before-present, with present = 1950 CE) atmospheric CH₄, as reconstructed from ice cores, nearly doubles doubled from ~380 ppb at LGM to 695 ppb at 10 ka BP (Köhler et al., 2017), with very rapid concentration changes of about 150 ppb occurring during the transitions from the Older Dryas into the Bølling Allerød (BA), from the BA into the Younger Dryas (YD), and from the YD into the Preboreal (PB) / early Holocene (Figure 1). Furthermore, while Holocene atmospheric CH₄ is was very similar for 10 ka BP and PI (694 ppb, mean concentration for 300 a BP to 200 a BP), CH₄ decreases decreased linearly by 15% at from 10 to 5 ka BP and increases increased again linearly towards PI. If we assume that the atmospheric lifetime of CH₄ did not change dramatically between the LGM and the present, these changes in atmospheric CH₄ would require large changes in CH₄ emissions.

The change in methane between LGM and PI has been investigated in a number of studies. Some have used box models to explain the methane changes observed in ice cores. Recently Bock et al. (2017), for example, pointed to tropical wetlands as the main driver of glacial-interglacial CH₄ change from a study of methane isotopes from ice cores. In addition there are studies with comprehensive models. Kaplan (2002) investigated wetland CH₄ emissions during the LGM and the present

using the BIOME4 model. He finds wetland emissions of $140 TgCH_4yr^{-1}$ ($1Tg = 10^{12}g$) for the present-day situation and $107 TgCH_4yr^{-1}$ (-24%) for the LGM, with wetland areas at the LGM slightly larger than at present. Valdes et al. (2005) performed time-slice experiments with the Hadley Centre coupled model (HadCM3), using the Sheffield Dynamic Global Vegetation Model (SDGVM) as a fire and wetland methane emission model, as well as an atmospheric chemistry model. They find PI wetland CH₄ emissions of $148 TgCH_4yr^{-1}$ and LGM emissions of $108 TgCH_4yr^{-1}$ (-27%), with tropical sources changing rather little and NH high latitudes contributing most of the change in emissions. Emissions from biomass burning change from $11 TgCH_4yr^{-1}$ at PI to $7 TgCH_4yr^{-1}$ at LGM (-36%), contributing to the total emission change from $199 TgCH_4yr^{-1}$ at PI to $152 TgCH_4yr^{-1}$ at LGM (-24%). Weber et al. (2010) investigated wetland emissions for PI and LGM time slices with climate forcings from the Paleo Model Intercomparison Project PMIP2 ensemble, applied to an offline wetland CH₄ model. They found an overall reduction by 29-42%, with sources in the NH extratropics reduced by 51-60%, while tropical sources were reduced by 22-36%. Finally Hopcroft et al. (2017) investigated methane emission changes between LGM and PI using the Hadley Centre Global Environmental Model (HadGEM2-ES), considering wetlands, termites, biomass burning as CH₄ sources, along with ocean and geological emissions. They obtain an overall source reduction by 28-42%, with LGM wetland emissions (97 $TgCH_4yr^{-1}$, 80 $TgCH_4yr^{-1}$ if northern peatlands considered explicitly) reduced by 30% in comparison to PI (138 $TgCH_4yr^{-1}$), and termite emissions reduced by 40%.

Studies of time slices between the LGM and PI are much sparser. Kaplan et al. (2006), using BIOME4-TG as a terrestrial methane emission model also determining emissions of biogenic volatile organic compounds (BVOCs), and an atmospheric chemistry model, investigated time slices every 1000 years from LGM to the present. They found that changes in atmospheric CH₄ are largely due to a changed lifetime, mainly through BVOC emission changes. Interestingly, they find a larger wetland area for the LGM than for present-day, with emissions roughly the same ($\sim 110 T_g CH_4 yr^{-1}$), and an emission maximum around 10 ka BP. Finally, Singarayer et al. (2011) investigated methane for 65 time slices between 130 ka BP and PI with HadCM3 and SDGVM as a methane emission model. They point to orbital changes driving the methane increase between 5 ka BP and PI, as insolation increases in the SH tropics.

What these studies have in common is that they require (in some cases substantial) changes in the atmospheric lifetime of methane to explain the changes in atmospheric CH₄ reconstructed from ice cores. However, Levine et al. (2011) found very small changes in CH₄ lifetime between LGM and PI using the TOMCAT (Toulouse Off-line Model of Chemistry And Transport) atmospheric chemistry model, and Gromov et al. (2019), investigating which has also been found by Murray et al. (2014) and Hopcroft et al. (2017), who investigated CH₄ lifetime at the LGM using the EMAC model (ECHAM/MESSy Atmospheric Chemistry), also find a very similar lifetime. Therefore using different atmospheric chemistry models. This small change in CH₄ lifetime arises from the balance of several competing influences, such as changes in reactive compounds, changes in reaction rates due to temperature changes, and changes in atmospheric OH due to water vapour changes. As a result, substantial changes in emissions are required to explain the changes in atmospheric methane as seen in ice cores, and none of the previous studies have managed to obtain the required changes in methane emissions, while fulfilling the constraints of the present-day methane budget at the same time.

In the present-day top-down CH₄ budget (Saunois et al., 2016), 59% of the emissions are from anthropogenic sources and can therefore be ignored for times before a significant human impact on the methane budgetpre-anthropogenic times. However, 41% (231 TgCH₄yr⁻¹) of the emissions are from natural sources and are therefore relevant for the entire time since the LGM. In the top-down budget, 167 TgCH₄yr⁻¹ (72% of the natural emissions) are emitted from natural wetlands, and 64 TgCH₄yr⁻¹ come from "other" sources. These are not differentiated further in the top-down budget, but the bottom-up budget lists freshwater bodies (lakes), geological sources, wild animals, wildfires, permafrost soils and vegetation as further onshore (land) sources and geological and "other" as offshore (oceanic) sources.

We aim to assess the changes in the natural sources of methane from the LGM to the present in order to determine the factors driving the changes in atmospheric CH₄. We use a methane-enabled version of MPI-ESM, the Max Planck Institute Earth System Model, to investigate changes in natural methane emissions for six time slices from the LGM to the present. In this model we include submodels for methane fluxes from wetlands, termites and wildfires, but of the other natural methane fluxes listed above, many cannot easily be derived from the climate model state and therefore are neglected for now. As time slice experiments very likely are not helpful for looking into the BA-YD and YD-PB transitions, we neglect these for now, focusing instead on the longer timescale changes in methane.

2 Model and experiments

75 2.1 MPI-ESM 1.2

We use the Max Planck Institute Earth System Model (MPI-ESM) in version 1.2 (Mauritsen et al., 2019), the version to be used in CMIP6. All experiments are performed in resolution T31GR30 (Mikolajewicz et al., 2018). In comparison to the CMIP5 version (Giorgetta et al., 2013), a number of errors were corrected in the atmosphere and ocean models, and the land surface scheme JSBACH (Reick et al., 2013; Brovkin et al., 2013; Schneck et al., 2013) has been updated with a multilayer hydrology scheme (Hagemann and Stacke, 2015), the SPITFIRE fire model (Thonicke et al., 2010; Lasslop et al., 2014), and the improved soil carbon model YASSO (Tuomi et al., 2009; Goll et al., 2015).

2.2 Wetland methane emission model

The present-day area that wetland methane emissions originate from is highly uncertain. The generation of methane in the soil is dependent on plant composition, carbon content and carbon quality, essentially ecosystem properties, as well as the degree of anoxia in the soil, which depends on soil structure and water content, essentially hydrological properties. As there is no better estimate of the methane-generating area available, we determine the surface inundation and assume that this is a useful approximation of the areas where methane is generated.

2.2.1 Dynamic inundation model

We use an approach based on the TOPMODEL hydrological framework (Beven and Kirkby, 1979) to determine inundation extent dynamically. TOPMODEL is a conceptual rainfall-runoff model, based on the compound topographic index (CTI) $\chi_i = \ln(\alpha_i/\tan(\beta_i))$ with α_i a dimensionless index for the area draining through point i and β_i the local slope at that point. TOPMODEL determines the local water table z_i in point i in relation to the grid cell mean water table \bar{z} :

$$z_i = \bar{z} + \frac{1}{f} \left(\chi_i - \bar{\chi} \right) \tag{1}$$

with χ_i the local CTI index in point i, $\bar{\chi}$ the grid cell mean CTI index, and f a parameter describing the exponential decline of transmissivity with depth. From Eq. 1 we determine the grid cell fraction with a local water table depth $z_i \geq 0$. Since inundated areas become unreasonably large in some locations, we limit the valid range of CTI values by introducing the constraint $\chi_i \geq \chi_{min}$ following Stocker et al. (2014), with χ_{min} constant in space and time. We assume this to be the inundated and therefore methane-emitting area A_{inun} , unless soils are frozen. In these cases we determine the fraction of liquid water in the soil f_{liq} from the soil temperature T_{soil}

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$$f_{liq} = \begin{cases} 1 & \forall T_{soil} > 273.65K \\ 0.1 & \forall T_{soil} < 272.75K \\ (T_{soil} - 273.65K)K^{-1} & otherwise \end{cases}$$

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(limiting f_{liq} to $0.1 \le f_{liq} \le 1$ for numerical reasons), and determine the inundated area as $A'_{inun} = A_{inun} \times f_{liq}$, reducing the inundated area under freezing conditions, as frozen soils emit less methane, similar to Gedney and Cox (2003).

To determine the grid cell mean water table position \bar{z} , we determine the layer saturation $\Psi_k = \Theta_k/\Theta_{fc}$ for each soil layer k by dividing the volumetric moisture content Θ_k by the field capacity Θ_{fc} . Starting from the bottom of the soil column, \bar{z} is located in the first soil layer l with layer saturation Ψ_l less than the saturation threshold Ψ_{thres} . The final water table position then is

$$\bar{z} = z_{h,l} - \Psi_l \Delta z_l \tag{2}$$

with $z_{b,l}$ the bottom of soil layer l, and Δz_l the height of soil layer l.

Values for f, χ_{min} , and Ψ_{thres} were determined from sensitivity experiments. In the experiments described here, we use f=2.6, $\chi_{min}=8.5$, and $\Psi_{thres}=0.95$. Furthermore, comparison with remote sensing data (Prigent et al., 2012) showed that inundated area in grid cells with a mean CTI index $\bar{\chi} \leq 5.5$ is negligible. Inundation is therefore only determined for grid cells with $\bar{\chi} > 5.5$.

We use the CTI index product by Marthews et al. (2015) for the CTI index at a resolution of 15" in all present-day land areas, while we determine CTI index values for shelf areas that are below sea level at present, but above sea level under glacial conditions, from the ETOPO1 dataset (Amante and Eakins, 2009) using the topmodel library for R (Buytaert, 2011). In order to reduce storage requirements, we approximate the distribution of CTI values within a model grid cell by fitting a gamma distribution, following Sivapalan et al. (1987).

2.2.2 Wetland methane production and transport

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We use the methane transport model by Riley et al. (2011) to determine wetland methane emissions, with minor modifications to adapt the model to the vegetation and carbon cycle representation in JSBACH. The Riley et al. (2011) model determines CO₂ and CH₄ production in the soil, transport of CO₂, CH₄ and O₂ through the three pathways diffusion, ebullition and plant aerenchyma, as well as the oxidation of methane during transport.

Adaptations are described in the following. In the grid cell fraction determined to be inundated by the inundation model, soil organic matter (SOM) is decomposed under anaerobic conditions in the YASSO soil carbon model (Goll et al., 2015), assuming a reduction of decomposition by a factor of 0.35 (Wania et al., 2010) in comparison to the aerobic case. As YASSO is a zero-dimensional representation of soil C processes, we distribute the decomposition product to the soil layers according to the root distribution from Jackson et al. (1996). Partitioning of the anaerobic decomposition product into CO_2 and CH_4 is temperature-dependent, as in the original Riley model, with a baseline fraction of CH_4 production $fCH_4 = 0.35$ and a Q_{10} factor for fCH_4 of $Q_{10} = 1.8$ with a reference temperature of 295K.

For each grid cell, the methane model determines CH₄ production and transport for two grid cell fractions, the aerobic (non-inundated) and the anaerobic (inundated) fraction of the grid cell. If the inundated fraction changes, the amounts of CO₂, CH₄ and O₂ are conserved, transferring gases from the shrinking fraction to the growing fraction, proportional to the area change. While vegetation in JSBACH is determined for vegetation tiles, allowing a fractional coverage of plant functional types, the relevant properties (root distribution, SOM decomposition) are aggregated to grid cell level for the methane transport model for performance reasons. Previous sensitivity experiments showed that differences to a tile-resolving formulation are negligible.

2.3 Methane emissions from wildfires

To determine methane emissions from wildfires, we use the biomass burned, diagnosed from the SPITFIRE fire module (Thonicke et al., 2010; Lasslop et al., 2014), as well as information on vegetation composition from the dynamical vegetation model. We use the methane emission factors from Kaiser et al. (2012), mapped to the JSBACH plant functional types, to determine the fraction of burned biomass emitted as methane. Therefore changes in fire-related methane emissions are completely determined by changes in fire carbon emissions. Fire occurrence in the SPITFIRE model is determined as a function of flammability (higher under dryer/warmer conditions) and ignition probability, with ignition probability a function of lightning frequency and population density. We are currently limited to a fixed lightning distribution reflecting modern conditions, and we are assuming a population density of zero for all time slices earlier than preindustrial. Therefore the main factors affecting fire-related methane emissions are carbon content and moisture conditions.

For PI and PD we use an estimate of population density to determine the ignition probability, with ignition probability increasing with population density. However under very high population densities it is assumed that fire suppression increases, thus decreasing fire probability, and thus fire methane emissions, for very high population densities.

2.4 Methane emissions from termites

150 Methane emissions from termites are determined following the approach developed by Kirschke et al. (2013) and elaborated by Saunois et al. (2016), adapted for the use in a dynamical vegetation model. They distinguish between termite emissions from tropical and non-tropical areas, using different parameterisations for determining the termite biomass $M_{termite}$ and different emission factors for the two areas. For tropical areas, in our case defined as areas covered by the plant functional types (PFT) tropical broadleaf evergreen tree, tropical broadleaf deciduous tree, and C4 grass, we determine $M_{termite}$ from the annual gross 155 primary production GPP using $M_{termite} = 1.21 \times \exp(GPP \times 0.0008)$ (Kirschke et al., 2013). From $M_{termite}$ we determine methane emissions using an emission factor of $2.8 \mu gCH_4/gTermite/h$ (Saunois et al., 2016). The non-tropical areas, i.e. the areas where the tropical PFTs do not occur, with termite emissions we define as the areas suitable for temperate broadleaf evergreen trees using bioclimatic limits from Sitch et al. (2003): temperature of the coldest month $T_c > 3$ °C and a growingdegree-day sum on the basis of 5°C $GDD_5 > 1200$ °C. In these areas we assume a constant termite biomass $M_{termite} = 3 g/m^2$ 160 and an emission factor of $1.7 \mu gCH_4/gTermite/h$ (Saunois et al., 2016). If croplands occur in any particular grid cell (not relevant for experiments presented here), emissions from the cropland tile are reduced to 40% of the non-cropland grid cell mean emissions, also following Saunois et al. (2016).

2.5 Model experiments

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We performed model experiments for five time slices at 20 ka BP, 15 ka BP, 10 ka BP, 5 ka BP, and PI, in this case defined as the year 1850 CE. In addition we performed one transient historical experiment for the years 1850-2010 CE, starting from the PI time slice, in order to obtain a present-day (PD) climate state for evaluation purposes. All model experiments use prescribed orbital forcing from Berger (1978) and greenhouse gas forcings from Köhler et al. (2017). Orbital parameters and greenhouse gas concentrations are supplied to the model as 10 year mean vales and are updated every 10 model years. Atmospheric aerosols were constant at 1850 conditions (Kinne et al., 2013), with the exception of the historical experiment, and we considered no anthropogenic land use.

The time slice experiments were initialised from a – so far unpublished – transient model experiment from 26 ka BP to PI with prescribed ice sheet extent from the GLAC-1D ice sheet reconstruction (Tarasov et al., 2012; Briggs et al., 2014; Ivanovic et al., 2016). This model experiment was initiated at 26 ka BP and run transiently from then to PI, i.e. the year 1850 CE. Ice sheet extent, as well as bathymetry and topography (Meccia and Mikolajewicz, 2018) and river routing (Riddick et al., 2018) were continuously updated throughout the deglaciation.

As the original transient experiments did not contain the methane code required for the experiments described here, the time slice experiments were initialised from the transient experiment with a three-step procedure to minimise climate drift from the original experiment. In the first step, all model components were initialised from the transient experiment, with the exception of the inundation and the methane model, which were initialised from scratch. The model was integrated for 20 years from this initial state. This was repeated for a second time, but using the inundation and methane states reached at the end of the initial experiment. In a third step, the model was run for forty years, using the inundation and methane state reached at the end of

step two, while using the conditions of the transient model experiments for all other model components. In this way we insured ensured that the state of the physical model, as well as the biogeochemistry, would always be as close as possible to the model state in the fully transient experiment.

Present-day (PD) conditions we assess by performing a historical experiment for 1850-2010, initialised from the PI state of the transient deglaciation experiment. In the PD experiment we change GHG and atmospheric aerosol transiently, using the Stevens et al. (2017) aerosol parameterisation, but we do not consider anthropogenic land use.

In order to further assess the changes in methane emissions between PI and 20 ka BP, we have performed additional sensitivity experiments, starting from the PI state. In experiment PI-C-LGM we impose 20 ka BP land carbon stocks in all grid points that are common between 20 ka BP and PI, and in experiment PI-CO2-LGM we impose 20 ka BP atmospheric CO_2 on the land biogeochemistry, but not on the atmospheric physics. These experiments allow us to assess the effects of soil carbon and CO_2 changes on CH_4 emissions, while keeping climate unchanged. In experiments PI-noQ10 and 20k-noQ10, finally, we modified the CH_4 production parameters ($fCH_4 = 0.28$ and $Q_{10} = 1.0$) in such a way that partitioning between CO_2 and CH_4 is no longer temperature dependent, while giving similar total emissions in the PI state, thus mimicking the behaviour of models that do not take this factor into account.

Climate in the preindustrial state is very similar to the preindustrial control experiment described in Mauritsen et al. (2019) and Mikolajewicz et al. (2018). However, the orography used in the present experiments is different from that in the published preindustrial control experiments. The latter experiments employ a mean orography, while the orography in the transient deglaciation experiment that we used as starting conditions for our time slice experiments, is an envelope orography. In the envelope orography, the grid-cell elevation is enhanced in comparison to the mean orography, in order to better represent the influence of topography on atmospheric circulation.

For all experiments we analyse a 30 year mean climatology, with the exception of the PD experiment, where we analyse a 10 year mean climatology obtained from years 2000-2009. All plots of absolute emissions are shown on the land-sea mask appropriate for the time interval under consideration, while difference plots are shown on show the outline of the PI land-sea mask.

3 Results and discussion

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3.1 Evaluation of present-day methane emissions

3.1.1 Surface inundation

For the assessment of wetland methane emissions, the wetland area can to some extent be measured directly from satellites.

Remote-sensing products of surface inundation are available, for example by Prigent et al. (2012) and Schroeder et al. (2015).

To assess the quality of the modelled surface inundation, we rely on the Prigent et al. (2012) data set. However, four points need to be considered when comparing these data to model results:

1. The remote-sensing process is unable to penetrate snow cover, so snow-covered areas are considered non-inundated.

2. The remote-sensing product shows all inundated areas, including areas flooded as a result of anthropogenic processes, such as the creation of reservoirs and rice-paddies, which are not considered in the model.

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- 3. Remote-sensing may be unable to penetrate dense forest canopy, implying that inundation estimates may be biased in forested areas.
- 4. Not all methane-generating areas have water tables above the surface. Water tables in northern peatlands, for example, tend to be below the surface for part of the year, especially in the summer.
- In order to make model output and remote-sensing data comparable, we therefore mask all snow-covered areas in the model output, and we use data on rice-growing areas by Monfreda et al. (2008) to mask all rice-growing areas from both the remote-sensing data and the model output. After these modifications, modelled inundated areas for the present-day period (mean over 2000-2009) are slightly larger than those observed by Prigent et al. (2012) (mean over 1993-2007) (Fig. 2). For the tropics (TRO, here for simplicity defined as latitudes between 30°N and 30°S) the annual mean inundated area is $1.2 \times 10^6 km^2$ in the model results, while Prigent et al. show $0.8 \times 10^6 km^2$. The seasonality is phase-shifted, with the model showing the peak inundation in April, while Prigent et al. show the inundation peak in August (Fig. 2). For the glacier-free NH extratropics (NXT, here defined as north of 30°N), the seasonality of inundation is similar in observations and model, but the summer peak in inundation is larger in the model ($2.5 \times 10^6 km^2$ for the JJA mean) than in the observations ($2.3 \times 10^6 km^2$).

Comparing the spatial pattern of the annual maximum inundation (Fig. 3), the overall pattern is rather similar, although two major differences are apparent: 1) The annual maximum inundation is more localised in the observations, while it is less clearly defined and reaching lower maximum values in the model, and 2) after removal of the rice-growing areas the model does not show a significant inundation maximum in India, very. These differences are likely due to the low resolution of the model we are employing. At higher resolutions, inundation becomes more localised, thus (1) appears clearly related to resolution, while (2) is due to an underestimate of the Indian monsoon precipitationin, also likely due to the low model resolution. We thus judge the methane generating areas produced by the model as reasonable, keeping in mind the likely low bias of the remote-sensing inundation product.

As described above, we use the inundated area to determine the methane emitting area. To evaluate the inundated areas leading to the wetland emissions, it has to be kept in mind that NXT emissions mainly are from the summer season, implying that the JJA (June, July, August) mean inundation is relevant for these, while the seasonality of TRO emissions is much less pronounced, implying that the annual mean inundation is relevant. In the following we therefore assess the effective inundated area, defined as the annual mean inundated area in tropical latitudes (TRO, between 30°N and 30°S), and the JJA (June, July, August) mean inundated area in the glacier-free NH extratropics (NXT, north of 30°N). For the present-day climate state, the effective inundated area is $1.5 \times 10^6 km^2$ in TRO and $2.6 \times 10^6 km^2$ in NXT (differences to the numbers shown above due to the removal of rice-growing areas in the comparison to observations).

245 3.1.2 Natural methane emissions

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So far it has not been possible to directly measure the quantity – surface methane fluxes – that we aim to assess in this publication on appropriate scales. Methane flux measurements exist for single sites of meter scale, mainly using measurement chambers, and for slightly larger scales, using eddy-covariance towers, but so far the scales relevant for global scale modelling, the model grid-cell to global scales, have not been covered by direct methane flux measurements (Melton et al., 2013; Saunois et al., 2016; Poulter et al., 2017). For assessment of our model experiments we therefore need to rely on global assessments (Saunois et al., 2016), and we can gain some additional insight from atmospheric inversions (Bousquet et al., 2011).

Under present-day (PD) climatic conditions (i.e. years 2000-2009 in the transient historical experiment), the model simulates wetland methane emissions of $222TgCH_4yr^{-1}$ (209 – $239TgCH_4yr^{-1}$), fire emissions of $17.6TgCH_4yr^{-1}$ (15.6 – $18.8TgCH_4yr^{-1}$), termite emissions of $11.7TgCH_4yr^{-1}$ (10.8 – $12.2TgCH_4yr^{-1}$), and a soil uptake of $17.5TgCH_4yr^{-1}$ (17.4 – $17.7TgCH_4yr^{-1}$). The values shown are mean values over the years 2000-2009 of the historical experiment, with the value in brackets giving the minimum and maximum annual emissions occurring in the model results. These values fall well within the ranges reported by Saunois et al. (2016), who report $153 - 227TgCH_4yr^{-1}$ for natural wetlands, $27 - 35TgCH_4yr^{-1}$ for biomass and biofuel burning, with biofuel burning making up 30-50%, $3 - 15TgCH_4yr^{-1}$ for termites, and $9 - 47TgCH_4yr^{-1}$ for the soil uptake. Spatial patterns of modern emissions (Fig. A2 and A3) are generally similar to those shown by Saunois et al. (2016).

Furthermore, wetland methane emission estimates from atmospheric inversions (Bousquet et al., 2011) show that the majority (62-77%) of the PD emissions come from the TRO region, while a much smaller part (20-33%) are emitted from NXT. Of the modelled total wetland CH₄ emissions for PD conditions, $156TgCH_4yr^{-1}$ (70%) are from TRO and $65TgCH_4yr^{-1}$ (29%) are from NXT, while emissions from the SH extratropics (here defined as south of 30°S) are negligible. The latitudinal distribution of modelled PD wetland methane emissions therefore is well within the range obtained from atmospheric inversions.

Overall, the PD state is rather similar to the PI state assessed in the following section, with very small differences in the distribution of emissions (Figures A2 and A3), but generally higher methane emissions. At 287.4 K, the global mean temperature in the PD climate state is 0.5 K warmer than preindustrial (Table 1). Precipitation is similar, leading to negligible differences in the effective inundation. With 1140 PgC, 8% larger than PI, the global soil C stock is also rather similar. However, vegetation productivity is enhanced in comparison to PI, due to warmer temperatures and higher CO₂ concentrations. The net methane emissions in PD climate are 29% larger than PI (Table 2), with wetland methane emissions 33% larger, with a larger increase (+42%) in TRO than in NXT (+16%). Fire emissions are 18% larger than PI, termite emissions increase by 66%, and the soil uptake increases by 140%. The latter increase is largely due to the higher atmospheric concentration of CH₄, which drives additional methane into the soils in comparison to the lower-CH₄ PI state. The larger fire emissions are mainly due to higher population densities in the 2000s than in 1850, although the very high population densities in eastern North America, Europe, and southern Asia are assumed to drive an increase in fire suppression in the SPITFIRE model (Lasslop et al., 2014). Thus fire emissions are decreased here, despite the general increase in most other places. Termite emissions are higher in the modern

climate due to an increase in GPP under higher CO_2 , while wetland emissions largely increase due to the higher temperatures of the modern climate, with CO_2 -fertilisation playing an additional role.

3.2 Preindustrial methane emissions

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The climate in our PI experiment is very similar to the one described by Mikolajewicz et al. (2018). The global mean near-surface air temperature is 286.9 K (Table 1). The annual mean temperature in the TRO area T_{TRO} is 294.5 K, while T_{NXT} , the annual mean temperature in the NXT area, is 275.2 K. The NH ice sheet area is limited to Greenland, with the ice sheet having an area of $1.8 \times 10^6 km^2$ in our model setup. Under these climatic boundary conditions, we obtain total net methane emissions of $181 TgCH_4 yr^{-1}$ (Table 2), with wetlands contributing $167 TgCH_4 yr^{-1}$ (Fig. 4a), fire and termites $15 TgCH_4 yr^{-1}$ and $7.0 TgCH_4 yr^{-1}$, respectively (Fig. 5a and b), while the soil uptake is $7.3 TgCH_4 yr^{-1}$ (Fig. 5c).

Wetland emissions, the dominant natural component of the terrestrial methane fluxes, mainly originate in TRO ($110 TgCH_4 yr^{-1}$), while emissions from NXT are $56 TgCH_4 yr^{-1}$ (Table 2). The main factors determining wetland methane fluxes, apart from temperature, are the emitting area and the soil carbon stock that the soil respiration (and thus methane production) is derived from. In the PI state the global effective inundated area is $4.0 \times 10^6 km^2$ (Fig. 4b), of which $2.7 \times 10^6 km^2$ are located in NXT, while $1.3 \times 10^6 km^2$ are in TRO (Table 1). For soil carbon, on the other hand, the global stock is 1054 PgC ($1Pg = 10^{15} g$), with most of the soil carbon (588 PgC) located in NXT, while it is 439 PgC in TRO.

Methane emissions from fires (Fig. 5a) closely follow the fire distribution, with most fire methane emissions coming from subtropical Africa and South America, although some emissions also originate in North America, Southern Europe and South Asia. Termite emissions, on the other hand mainly originate from tropical regions, especially southern Asia, with minor contributions from subtropical regions on all continents (Fig. 5b). Methane uptake by upland soils (Fig. 5c), finally, is distributed widely with no large regional variations.

3.3 Wetland methane emissions

Under LGM boundary conditions the global mean temperature is 4.4 *K* colder than under PI conditions (Table 1). Extensive glaciers cover the NH extratropics and sea level is lower, leading to a 15% increase in total land area, although total glacier-free area is nearly identical (Table 1). The TRO area A_{TRO} thus is 12% larger than PI, while the NXT area A_{NXT} (by definition glacier-free) is 14% smaller. The temperature decrease is less pronounced in TRO (-3.1 *K*) than in NXT (-5.8 *K*). Precipitation decreases by 10% in the global mean, with an 11% decrease in TRO and a 19% decrease in NXT. TRO effective inundated area I_{TRO} thus increases by 19% (Table 1, Figure 6 a, Figure A4 a), while NXT effective inundated area I_{NXT} decreases by 6%. The global soil C stock is 617 *PgC*, substantially smaller (-41%) than at PI, with the decrease smaller in TRO (-33%) than in NXT (-49%). As a result of these climate changes, wetland methane emissions decrease by 51% (Table 2, Figure 6 e, Figure A4 e), with a TRO emission decrease of 47%, while NXT emissions decrease by 59%, with the majority of the latter emissions coming from areas in East Asia adjacent to the Yellow Sea and North America south of the Laurentide ice sheet. The wetland CH₄ emissions therefore decrease nearly everywhere (Figure 6 e), with one major exception: The continental shelf areas that

are exposed due to the lower sea level become significant sources of methane, especially in Indonesia, but also in Africa and Asia. At $29 T gCH_4 vr^{-1}$, they contribute about 35% of the total wetland CH₄ emissions at the LGM.

Analysing the factors leading to the reduction in 20 ka BP wetland CH₄ emissions further (see Appendix A for details), we can explain a reduction of emissions by 42% in comparison to PI by the changes in land carbon and atmospheric CO₂. Changes in temperature lead to a further reduction by about 20%, while changes in available land points actually lead to an *increase* in emissions, as the available land points in the northern high latitudes (with low emissions) are reduced, while the available land points in the tropics (with high emissions) are increased.

For 15 ka BP, the global mean temperature change is $-2.8\,K$, relative to PI. NH ice sheet extent is 24% lower than at LGM, but still extensive. The total land area is 12% larger than PI due to the lower sea level, but A_{nonglac} in only larger by 1%. A_{NXT} is thus reduced by 10% (Table 1), while A_{TRO} is increased by 11%. The change in T_{TRO} is $-2.0\,K$ (Table 1), while it is $-3.8\,K$ for T_{NXT} . Precipitation decreases by 6% in the global mean, with a 4% decrease in TRO and a 10% decrease in NXT. Precipitation in NH Africa is slightly increased due to a stronger West African monsoon. I_{TRO} thus is larger by 29%, while I_{NXT} is 8% smaller than PI (Table 1, Figure 6 b, Figure A4 b). Global soil C is at 815 PgC (-23%), with TRO C stocks at 398 PgC (-9%), while NXT stocks are at 392 PgC (-33%). Total wetland methane emissions decrease by -22% as a result, with TRO emissions decreasing by 17% and NXT emissions of by 33% (Table 2, Figure 6 f, Figure A4 f). In contrast to the LGM situation, there is an increase in CH₄ emissions from NH (sub-) tropical Africa Figure 6 f) to $19\,T_{g}CH_{4}\,yr^{-1}$ (+58%), due to wetter conditions in the Sahel area. The exposed shelf areas emit about $38\,T_{g}CH_{4}\,yr^{-1}$ overall, 29% of the total emissions.

For 10 ka BP, our model indicates a global mean temperature change of -0.7K (Table 1). Glacial area is much reduced in comparison to the LGM, but remains of the Laurentide ice sheet still cover parts of north-eastern Canada, leading to a lower sea level than at PI. The total land area thus is 3% larger than at PI (A_{nonglac} -1%), with A_{TRO} 3% larger due to lower sea level and A_{NXT} 4% smaller due to the remaining ice sheet coverage. The temperature decrease is larger in TRO (-1.0K) than in NXT (-0.6K). Precipitation is near PI levels in the global mean (-1%), with a 6% increase in TRO, and a 1% decrease in NXT. Precipitation in NH Africa is strongly increased due to a strong West African monsoon. I_{TRO} is increased by 35%, mainly indue to the wetter conditions in north Africa, while I_{NXT} is decreased by 12% in NXT (Table 1, Figure 6 c, Figure A4 c). Global soil C is at 983 PgC, quite near the PI total stock (-7%), with a TRO soil C stock of 449 PgC (+2%) and a NXT stock of 510 PgC (-13%). As a result, wetland CH₄ emissions are very similar to PI (Table 2), with +7% in TRO wetland emissions and -14% in NXT emissions (Table 2, Figure 6 g, Figure A4 g). The increase in TRO emissions mainly occurs in the Sahel area, where the West African monsoon is strongly increased, leading to more precipitation, increased inundated area, and more biomass and soil C. Emissions from NH Africa are $37TgCH_4yr^{-1}$ (+208%), an increase larger than the total increase in TRO emissions. Emissions from the (small) exposed shelf areas are at $8TgCH_4yr^{-1}$ (5% of the total wetland CH₄ emissions). NXT emissions are smaller than PI in North America and Europe, but they are larger than PI in northern Asia, due to the summer warming from the changed insolation at 10 ka BP.

At 5 ka BP, global mean temperature change is at -0.2K (Table 1). Ice sheet areas are as in the PI state, thus TRO and NXT areas are unchanged. T_{TRO} is slightly lower (-0.6K), but T_{NXT} is very similar to PI. Precipitation changes are very small in the global mean, with a a 4% increase in TRO and a 2% increase in NXT, with the West African monsoon slightly stronger than

PI. I_{TRO} increases by 15%, while I_{NXT} decreases by 1% (Table 1, Figure 6 d, Figure A4 d). Global soil C stocks are 1043 PgC, slightly smaller than preindustrial (-1%), with an increase by 4% in TRO, especially in the southern Sahel region, while NXT is 4% lower than PI. Total wetland methane emissions increase by 2%, with TRO and NXT wetland emissions both increasing by 2% (Table 2). Emissions from NH Africa are $21TgCH_4yr^{-1}$ (+75% compared to PI, 19% of TRO emissions), while emissions from the SH are generally decreased. NXT emissions are decreased in northern North America, while emissions from northern Asia and southern North America are increased.

3.4 Methane emissions from wildfires

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For all time slices before PI we assume that no humans were present, leading to a generally decreased probability of fire ignition in comparison to PI and PD. For the LGM (Fig. 7 a) fire CH₄ emissions (Table 2) are 73% smaller. As biomass is reduced strongly under the cold and low-CO₂ conditions of the LGM, fire-related C emissions are also reduced. At 15 ka BP (Fig. 7 b) fire emissions are 53% lower than PI, while they are 40% smaller at 10 ka BP (Fig. 7 c). Generally, the spatial pattern of emission changes at 15 and 10 ka BP mainly reflects precipitation changes: Enhanced emissions occur in areas where precipitation is reduced, enhancing vegetation flammability. In the Sahel area, this relationship is different, though. Here, the enhanced rainfall leads to an increase in vegetation cover, especially grass cover. As a result, more biomass is available for combustion, leading to enhanced emissions. At 5 ka BP, finally, fire emissions are reduced by 45%. As climate is already relatively similar to the PI situation, the main reason for the fire emission reduction here is the smaller ignition probability due to the absence of humans.

3.5 Methane emissions from termites

Termite emissions are mainly determined by gross primary productivity (GPP) in tropical and subtropical areas. The lower atmospheric CO₂ and temperature under LGM conditions decrease GPP everywhere. Therefore termite CH₄ emissions are reduced by 58% relative to the PI level (Table 2). For 15 ka BP, there also is a general reduction in termite emissions, with 4.6 TgCH₄ yr⁻¹ in total (-34%). However, the enhanced rainfall in the Sahel area leads to an increase in termite methane in this area. The latter is similar at 10 ka BP, where total emissions are 13% smaller in comparison to PI. The enhanced productivity in the Sahel therefore more than compensates the decrease in termite methane from the Amazon and African rain forests. At 5 ka BP, finally, termite emissions are slightly smaller than PI (-7%), with minor decreases in the rain forest areas and a slight increase in the Sahel.

3.6 Methane uptake by soils

The soil continually exchanges methane and oxygen with the atmosphere through diffusion. In areas where soil conditions are aerobic, methane concentrations in the soil are smaller than atmospheric concentrations, thus driving a flux of methane into the soil. In the soil the methane is oxidised, with oxidation rates dependent on the concentrations of CH_4 and O_2 , as well as

temperature. The gas exchange between soil and atmosphere is also modified by the presence of plants, as some plant tissues can transport gases between plant roots and leaves.

In our experiments, we find that the soil uptake of methane is to a large extent determined by the gradient of methane between soil and atmosphere. Thus higher atmospheric concentrations of methane directly lead to a larger soil uptake of methane. Under LGM conditions, the atmospheric CH₄ concentration is 370 ppb, slightly less than half the PI concentration. Consequently soil methane uptake decreases by 68% compared to PI (Table 2), with decreased temperatures being an additional factor (Fig. 9 a). At 15 ka BP (atmospheric CH₄ of 464 ppb), the soil uptake is 52% smaller, while it is changed by -14% at 10 ka BP (688 ppb) and -28% at 5 ka BP (579 ppb). Spatially, the change in methane uptake is rather uniform, showing a similar reduction in uptake in most locations (Fig. 9). The exception to this is, once again, the Sahel area, which shows an increase in methane uptake most pronounced for 10 ka BP (Fig. 9 c), but also for 5 ka BP (Fig. 9 d). For these time slices the increase in vegetation cover in the Sahel region leads to a localised increase in methane uptake.

3.7 Time slice comparison

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The net natural methane flux, i.e. the sum of all flux components, increases from $86 T g C H_4 y r^{-1}$ at 20 ka BP (-52% compared to PI) to $181 T g C H_4 y r^{-1}$ in the PI state, and $233 T g C H_4 y r^{-1}$ (+29%) at present.

The wetland emissions from TRO are the most important component of the net methane flux during all time slices. Their contribution is smallest at PI (61% of total net emissions) and largest at 20 ka BP (67%). The contribution from NXT ranges from 27% at 20 ka BP to 32% at 5 ka BP. Fire emissions make up 4-5% in the purely natural states between 20 ka BP and 5 ka BP, and about 8% for the anthropogenically influenced states at PI and PD. Termite emissions make up between 3.4 and 5.0% of net emissions, and soil uptake reduces the emissions by between 2.5% at 15 ka BP and 7.0% in the PD state (Table 2). Our modelled CH₄ emissions are also supported by the interpolar gradient in methane, as reconstructed by Baumgartner et al. (2012) for the LGM and Mitchell et al. (2013) for PI. Mitchell et al. infer tropical emissions of about $125TgCH_4yr^{-1}$ and NXT emissions of about $66TgCH_4yr^{-1}$ for the PI state (estimated from their Figure 1), while Baumgartner et al. infer tropical emissions of about $60TgCH_4yr^{-1}$ and NXT emissions of about $25TgCH_4yr^{-1}$ for the LGM (estimated from their Figure 5b).

In the modelled emissions, we are missing two components of the natural methane cycle: Wild animals and geological sources. For geological emissions, estimates vary widely, with bottom-up estimates in Saunois et al. (2016) of $35 - 76TgCH_4yr^{-1}$ for on- and offshore sources, while Petrenko et al. (2017), estimating methane ¹⁴C for the YD from ice cores, constrain methane stemming from old carbon reservoirs to the range $0 - 18.1TgCH_4yr^{-1}$ at the YD. These fluxes likely are constant in time, although there might be changes during periods of sea level rise and fall, as hypothesised for CO_2 by Huybers and Langmuir (2009). Methane emissions from wild animals, especially ruminants, are very difficult to estimate, current estimates for the present span a range $2 - 15TgCH_4yr^{-1}$ (Saunois et al., 2016), and estimates for other time slices are even less confident, but might be of the order of $15 - 20TgCH_4yr^{-1}$ for times before significant human influence (Chappellaz et al., 1993). In principle, these emissions should somehow be related to the net primary productivity, as this would determine the carrying capacity of the ecosystem, implying smaller fluxes in the glacial than in the Holocene. Adopting the ice-core based

estimate by Petrenko et al. (2017) for the geological fluxes, we can thus hypothesise these to be $9 \pm 9 T g C H_4 y r^{-1}$, while wild animals might add $15 \pm 10 T g C H_4 y r^{-1}$. The total unaccounted fluxes might therefore be of the order of $24 \pm 19 T g C H_4 y r^{-1}$.

To compare the net fluxes to the reconstructed atmospheric CH₄ concentrations from ice cores, we determined the implied methane emissions (Fig. 10). We converted the methane concentrations into a methane burden, using a conversion factor of $2.767TgCH_4ppb^{-1}$ (Dlugokencky et al., 1998). With a tropospheric lifetime of 9.3yrs (range given 7.1 - 10.6yrs), an approximation for the present-day situation (Saunois et al., 2016), we then determined the methane flux required to obtain the CH₄ concentration reconstructed for all time slices except the present-day period, which is dominated by anthropogenic emissions. From this flux we subtracted the unaccounted sources, as described above, to determine the implied emissions. Uncertainties from the unaccounted fluxes are represented as error bars, however the uncertainty in tropospheric lifetime is not considered here, but would roughly add another 15%.

Comparing the modelled net emissions to the implied emissions (Fig. 10), the modelled fluxes are within the range of uncertainty for all time slices except for 15 ka BP and 5 ka BP, with modelled net emissions larger than the implied fluxes for these time slices. The net emissions increase by more than 100% going from 20 ka BP to 10 ka BP and PI, we can thus explain the methane increase from LGM to Holocene with CH_4 emissions only, not requiring changes in methane lifetime. However, we so far cannot explain the Holocene changes in atmospheric CH_4 , decreasing between 10 ka BP and 5 ka BP, and increasing subsequently. For 5 ka BP, Singarayer et al. (2011) explain the decreased emissions in comparison to PI by reduced SH precipitation, leading to lower wetland CH_4 emissions. We obtain a similar result in our model, with SH wetland emissions reduced by $10T_3CH_4$ yr⁻¹ in comparison to PI. However our NH emissions are substantially larger than at PI, due to substantial increases in Sahel precipitation and wetland emissions. We assume that this is due to an overestimate of the West African monsoon and its impact on African methane emissions, as a general reduction in the West African monsoon would lead to decreases in TRO emissions for 15 ka BP, 10 ka BP, and 5 ka BP, bringing model results more in line with the implied emissions determined from ice core CH_4 . However, this is speculative at this point and would require further experiments.

4 Conclusions

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In this assessment we considered all natural emissions of methane, with the exception of emissions from wild animals and geological sources. In our experiments we found that it is possible to explain the difference between LGM (20 ka BP) and PI methane concentrations purely by changes in the emissions of methane, without requiring changes in the atmospheric lifetime of CH₄. The time slice experiments we performed suggest that there are three main drivers to changes in methane emissions over the time from the LGM to the present:

 Global mean temperature and CO₂: Higher atmospheric CO₂ concentrations increase NPP and thus also soil carbon available for anaerobic decomposition to CH₄. Similarly, higher global mean temperature also increases NPP and soil C decomposition, and it furthermore increases the ratio of CH₄ to CO₂ production in anaerobic decomposition. Thus, higher atmospheric CO₂ and higher global mean temperature lead to larger wetland emissions of CH₄. This affects emissions from wildfires and termites in a similar way, as fire C release is dependent on biomass and termite biomass is dependent on GPP and thus CO_2 and temperature.

2. Ice sheet area and sea level: Larger ice sheets remove CH₄ sources in the northern hemisphere extratropics as these are covered by the ice sheets, which is especially important for wetland methane emissions from North America. At the same time large ice sheets lower sea level, enlarging tropical wetland area as the continental shelf is exposed and becomes a significant source of methane. This is mainly relevant in the tropics, as high latitude shelf areas exposed under glacial conditions, for example the Laptev sea shelf, experience extremely cold conditions in glacial climate, leading to negligible methane emissions. Exposed shelf areas in Indonesia, Africa, and South America, on the other hand, emit significant amounts of methane. Thus lower sea level leads to larger emitting areas and thus higher emissions of methane.

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3. The West African monsoon: During the time slices when the West African monsoon is stronger than at present, i.e. at 15 ka BP, 10 ka BP, and 5 ka BP, precipitation in the Sahel region is significantly enhanced in comparison to the PI state, leading to an increase in vegetation cover, productivity and biomass burning. As a result, methane emissions at these times are stronger than at present, leading to a significant increase in (sub-)tropical CH₄ emissions, with all natural methane sources increased.

For methane emissions from wildfires, a further factor influencing the emissions is the human population density, as this strongly affects the fire probability in the SPITFIRE model employed in JSBACH (Fig. 10). The soil uptake of methane, on the other hand, is strongly dependent on the atmospheric concentration of methane (Fig. 10).

The changes in methane from LGM to the present are dominated by changes in tropical wetland emissions, with mid and high latitude wetland emissions being a significant but secondary factor, gaining in importance as the high latitudes become ice-free. In total the wetland emissions account for Wetland emissions make up 93 – 96% of the net CH₄ flux, and all other methane sources are thus of minor importance.

Code and data availability. The primary data, that is the model code for MPI-ESM, is freely available to the scientific community and can be accessed with a license on the MPI-M model distribution website. In addition, secondary data and scripts that may be useful in reproducing the authors' work are archived by the Max Planck Institute for Meteorology. They can be obtained by contacting the first author or publications@mpimet.mpg.de.

Appendix A: Overview over absolute emissions Sensitivity of LGM wetland fluxes

The large changes in wetland methane emissions between PI and 20 ka BP are due to a combination of several factors. While climate change is the ultimate cause of all of these influences, it is possible to identify several individual factors that likely contributed, and may have led to the larger changes in methane emissions than in some previous studies: (1) Due to sea level fall

and glacier expansion, the ice-free land domain changes appreciably, (2) reduced soil carbon at 20 ka BP leads to less organic matter being available to be decomposed, (3) reduced atmospheric CO₂ leads to lower vegetation productivity and litterfall, thus reducing both the plant transport of methane and the availability of soil organic matter for decomposition. In addition (4), the temperature-dependence of the ratio of CH₄ and CO₂ production in anaerobic soil organic matter decomposition is a relatively recent finding not considered in all methane transport model yet (Xu et al., 2016) and may also have played a role. In order to further assess these factors, we conducted a set of sensitivity experiments, as detailed in Section 2.5.

The ice-free land domain changes appreciably between PI and 20 ka BP (Fig. A1), with expanded ice sheets reducing the available area in the NH high latitudes, and lower sea level exposing additional land both in the high latitudes and in the tropics. Restricting analysis to the glacier-free land points common to both the PI and 20 ka BP time slices leads to a reduction in emissions by 16% for the PI state, and by 35% for the LGM state (Table 3). The change in ice sheets and sea level thus leads to an increase in 20 ka BP wetland emissions, compared to a state where the land-sea mask does not change, as some of the additional 20 ka BP grid points are located in the tropics, where large amounts of methane are emitted.

Imposing 20 ka BP land carbon stocks in experiment PI-C-LGM leads to a further reduction by 18% (evaluated for the common grid points), while imposing 20 ka BP atmospheric CO₂ for the land biogeochemistry in experiment PI-CO₂-LGM leads to a CH₄ emission reduction by 24 and 25%, evaluated for all grid points and the common grid points, respectively. The combined effect of 20 ka BP land carbon and CO₂ thus is a 42% reduction in emissions (common grid points). The remaining reduction by 20% of the PI emissions on the common grid points thus needs to to be explained by other factors, with temperature change being most important.

Another factor that may be different in our model in comparison to the models previously employed is the temperature dependence of the CO₂ to CH₄ partitioning during anaerobic decomposition. Disabling this temperature dependence in experiment 20k-noQ10 leads to a 16% increase in CH₄ emissions, with parameters chosen in such a way that the change is negligible (+3%) for the PI state. The enabled temperature dependence of the CO₂ to CH₄ partitioning in the standard setup thus is a factor that reduces 20ka BP emissions further.

Appendix B: Overview over absolute emissions

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For the convenience of readers who are more interested in absolute emissions than in the differences to the PI state, we also provide an overview over the absolute emissions for the present-day situation (Figs. A2 and A3), as well as the absolute wetland emissions for the past time slices (Fig. A4).

Author contributions. TK developed the methane module, performed model experiments and wrote the manuscript. UM performed the transient model experiment the climate states were derived from and contributed to climate model development. VB contributed to methane module development. TK and VB planned the study, and all authors discussed the analysis and the manuscript.

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

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Tables and Figures

Time	T _{GM}	T _{TRO}	T _{NXT}	A _{nonglac}	A _{TRO}	A _{NXT}	I _{TRO}	I _{NXT}
20 ka BP	282.5	291.5	269.4	133.9	72.3	54.3	1.5	2.5
15 ka BP	284.1	292.5	271.4	135.2	71.7	56.6	1.6	2.5
10 ka BP	286.2	293.6	274.6	133.2	66.1	60.6	1.7	2.4
5 ka BP	286.7	294.0	275.2	134.0	64.5	62.7	1.5	2.7
PI	286.9	294.6	275.2	134.1	64.5	63.0	1.3	2.7
present	287.4	295.1	276.0	134.1	64.5	63.0	1.5	2.6

Table 1. Climate and areas in experiments. Global mean annual temperature T_{GM} , TRO temperature T_{TRO} , and NXT temperature T_{TRO} , all in K. Global non-glaciated land area $A_{nonglac}$, TRO area A_{TRO} , NXT area A_{NXT} , TRO effective inundated area I_{TRO} , and NXT effective inundated area I_{NXT} , all in $10^6 km^2$.

Time	soil sink	wetland	fire	termite	net	TRO	NXT
20 ka BP	-2.3	81.8	4.0	3.0	86.4	58.1	23.3
15 ka BP	-3.5	129.6	7.0	4.7	137.7	91.5	37.7
10 ka BP	-6.2	165.8	8.8	6.1	174.5	117.3	48.3
5 ka BP	-5.2	170.7	8.1	6.5	180.1	112.6	57.7
PI	-7.3	166.7	14.9	7.1	181.4	110.1	56.3
present	-17.6	221.5	17.6	11.7	233.4	155.9	65.4

Table 2. Methane emissions for all time slices in $TgCH_4yr^{-1}$. Shown are soil uptake, total wetland emissions, fire and termite emissions, net emissions, and wetland emissions from TRO and NXT.

Experiment	CH ₄ , AP	$\Delta PI, \Delta P$	CH ₄ , CP	ΔPI, CP	∆, CP-AP
₽I	166.7		140.7		-16%
PI-C-LGM			116.0	-18%	
PI-CO2-LGM	127.1	-24%	105.4	-25%	
PI-noQ10	171.2	±3%	129.2	-8%	
20 ka BP	<u>81.8</u>	-51%	53.2	-62%	-35%
	·	∆20ka, AP		∆20ka, CP	
20k-noQ10	95.2	<u>+16%</u>	<u>67.4</u>	±27% ————————————————————————————————————	

Table 3. Sensitivity experiment results: wetland CH_4 emissions $[TgCH_4yr^{-1}]$ on all grid points (AP), emission change [%] with regard to PI state on all points, wetland emissions $[TgCH_4yr^{-1}]$ on the common grid points (CP), emission change [%] with regard to PI state on the common point, and emission change between common points and all points [%]. The reference state for emission change in experiment 20k-noQ10 is the 20ka BP state.

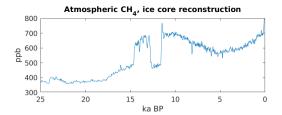


Figure 1. Atmospheric CH₄ as reconstructed from ice cores (Köhler et al., 2017) (Köhler et al., 2017, and references therein).

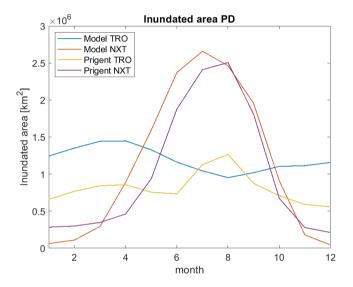


Figure 2. Climatology of monthly mean inundated area for model years 2000-2009 and Prigent et al. observations 1993-2007, separated for Tropics (TRO) and NH extratropics (NXT).

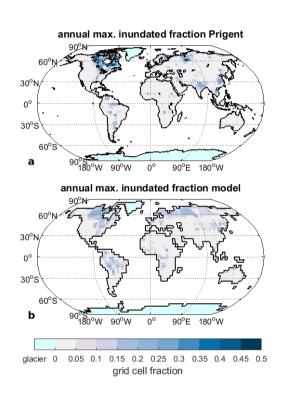


Figure 3. Annual maximum of mean monthly inundated fraction for Prigent et al. (a) and model (b).

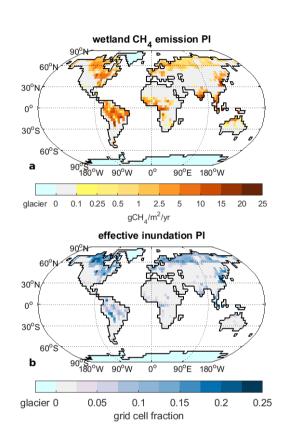


Figure 4. Wetland CH₄ emissions for preindustrial (PI) climate: Annual emissions of CH₄ from wetlands (a) and effective inundation (b).

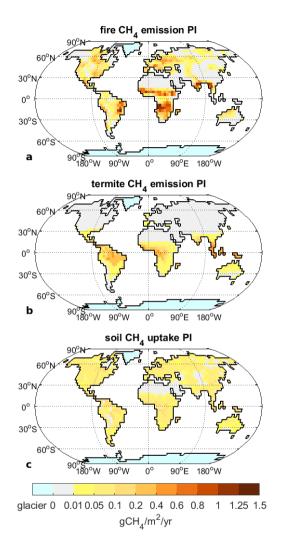


Figure 5. Non-wetland CH₄ emissions for PI climate: Annual emissions of CH₄ from fires (a) and termites (b), as well as annual soil uptake of CH₄ (c). In comparison to other fluxes, soil uptake has the opposite direction (sign).

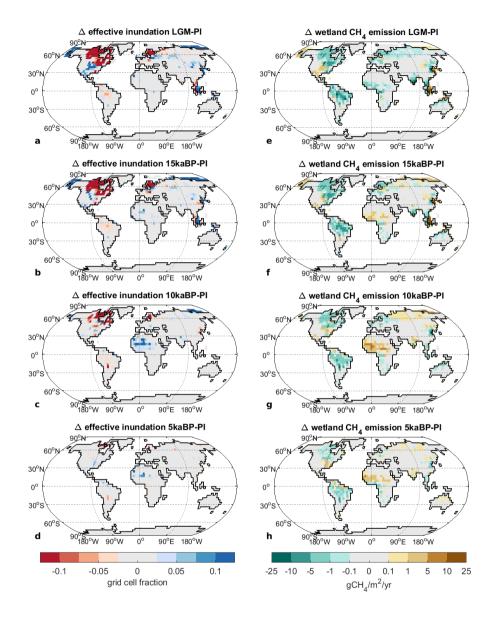


Figure 6. Change in effective inundation and wetland methane emissions for past climate states: a-d inundation difference to PI, e-h CH₄ emission difference to PI. a,e: LGM; b,f: 15 ka BP; c,g: 10 ka BP; d,h: 5 ka BP.

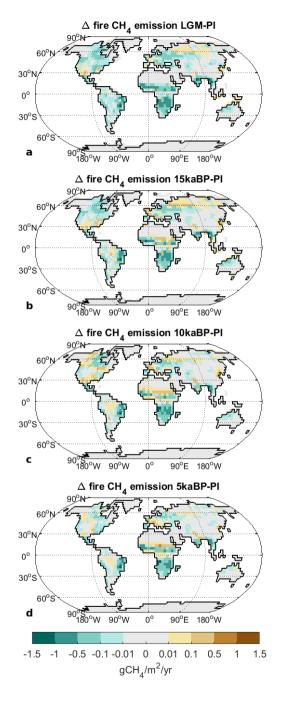


Figure 7. Difference in wildfire methane emissions to preindustrial for a) LGM, b) 15 ka BP, c) 10 ka BP, and d) 5 ka BP.

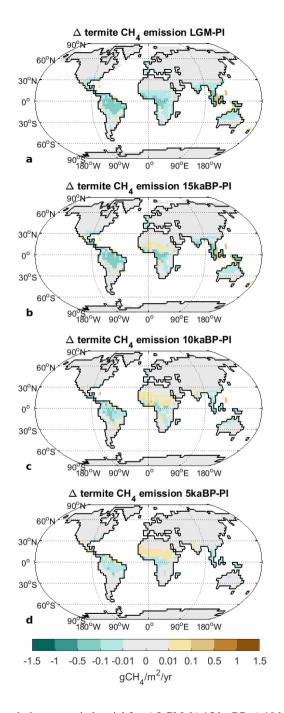


Figure 8. Difference in termite methane emissions to preindustrial for a) LGM, b) 15 ka BP, c) 10 ka BP, and d) 5 ka BP.

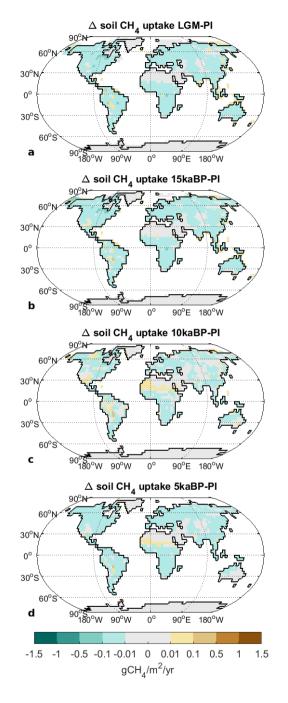


Figure 9. Difference in methane soil uptake to preindustrial for a) LGM, b) 15 ka BP, c) 10 ka BP, and d) 5 ka BP.

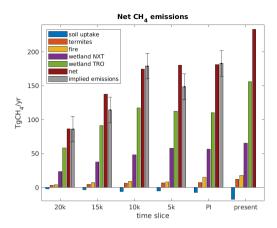


Figure 10. Components of the net CH_4 emissions for all timeslices. Soil uptake of CH_4 is shown as a negative flux. Calculation of implied emissions and error bar as detailed in the text.

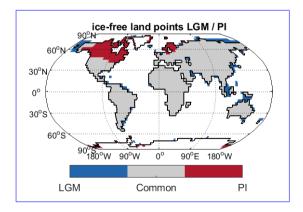


Figure A1. Ice-free land points for LGM and PI time slices, as well as grid points common to both.

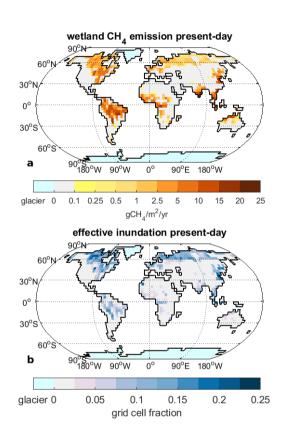


Figure A2. Wetland CH₄ emissions for present-day climate (2000-2009): Annual emissions of CH₄ from wetlands (a) and effective inundation (b). Please note the different colour scale for (b) in comparison to Fig. 3.

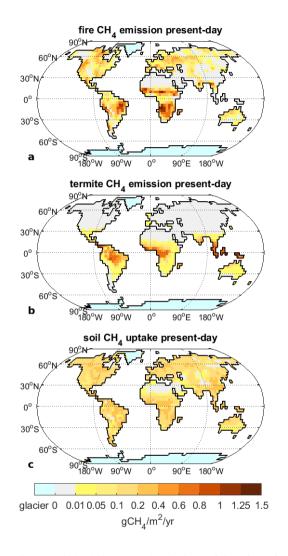


Figure A3. Model results for present-day climate (2000-2009): Annual emissions of CH₄ from fires (a) and termites (b), as well as annual soil uptake of CH₄ (c). In comparison to other fluxes, soil uptake has the opposite direction (sign).

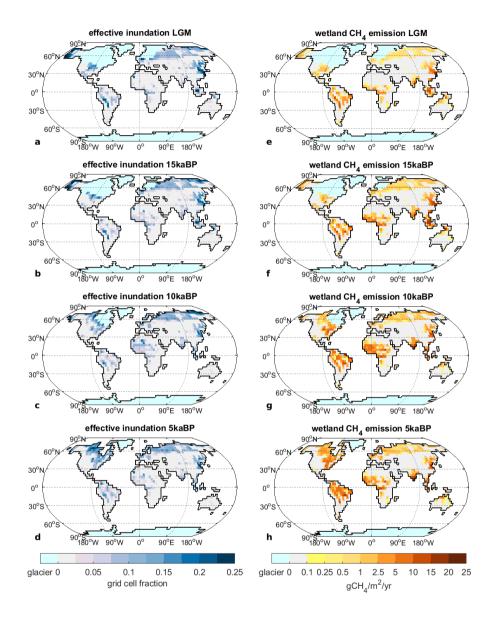


Figure A4. Absolute effective inundation and wetland CH₄ emissions for past climate states: a-d effective inundation, e-h wetland CH₄ emission. a,e: LGM; b,f: 15 ka BP; c,g: 10 ka BP; d,h: 5 ka BP.